

Patient-Reported Morbidity Instruments: A Systematic Review

Arvind Oemrawsingh, MD MHS^{1*}; Nishwant Swami, BA²; José M. Valderas, MD PhD MPH³; Jan A. Hazelzet, MD PhD¹; Andrea L. Pusic, MD MHS FACS FRCSC⁴; Richard E. Gliklich, MD⁵; Regan W. Bergmark, MD^{5, 6}

¹ Department of Public Health, Erasmus University Medical Center, Rotterdam, The Netherlands

² University of Massachusetts Medical School, Worcester, MA, USA

³ International Society for Quality of Life Research (ISOQOL), Health Services & Policy Research, University of Exeter Medical School, Exeter, England, UK

⁴ Division of Plastic and Reconstructive Surgery, Patient Reported Outcomes, Value, and Experience (PROVE) Center, Brigham and Women's Hospital, Boston, MA, USA

⁵ Department of Otolaryngology – Head and Neck Surgery, Harvard Medical School, Boston, MA, USA

⁶ Center for Surgery and Public Health, Patient Reported Outcomes, Value and Experience (PROVE) Center, Brigham and Women's Hospital, Boston, MA, USA

*Address correspondence to: Arvind Oemrawsingh, MD MHS

Room Na-2403, Department of Public Health

Erasmus Medical Center

P.O. Box 2040, 3000 CA

Rotterdam, the Netherlands

Abstract

Objectives

While (co)morbidities play an essential role in risk adjustment and outcomes measurement, there is little consensus regarding the best source of this data. The aim of this study was to identify (general) patient-reported morbidity instruments and their measurement properties.

Methods

A systematic review was conducted by using multiple electronic databases (Embase, Medline, Cochrane Central, and Web of Science) from inception to March 2018. Articles focusing primarily on the development and/or (subsequent) validation of a patient-reported morbidity instrument, were included. Following inclusion of relevant articles, measurement properties of each morbidity instrument was extracted by two investigators for narrative synthesis.

Results

A total of 1,005 articles were screened, of which 34 eligible articles were ultimately included. Most widely assessed instruments were the Self-Reported Charlson Comorbidity Index ($n = 7$), Self-Administered Comorbidity Questionnaire ($n = 3$) and Disease Burden Morbidity Assessment ($n = 3$). Most commonly included conditions were diabetes, hypertension and myocardial infarction. Studies demonstrated substantial variability in item-level reliability versus gold standard medical record review (κ range 0.66 – 0.86), meaning that the accuracy of self-reported comorbidity data is dependent on the selected morbidity.

Conclusions

The Self-Reported Charlson Comorbidity Index and the Self-Administered Comorbidity Questionnaire were the most frequently cited instruments. Significant variability was observed in reliability per comorbid condition of patient-reported morbidity questionnaires. Further research is needed to determine if patient-reported morbidity data should be used to bolster medical records data or serve as a stand-alone entity when risk adjusting observational outcomes data.

Keywords: Patient Report; Self Report; Surveys and Questionnaires; Psychometrics; Morbidity; Comorbidity; Health Services

Introduction

Value-Based Healthcare (VBHC) initiatives rely on risk adjustment to compare patient populations across hospitals. In addition to understanding the index disease of interest, comorbid conditions are necessary for case-mix adjustment. “Morbidity” is defined here as the presence of medical conditions. Clinicians are increasingly grappling with the challenges of treating patients with multiple co-occurring diseases (multi-morbidity). In addition to treatment difficulties, multi-morbidity is often associated with worse outcomes including decreased quality of life, psychological distress, longer hospital stay, more postoperative complications, higher cost of care, and higher mortality^{1,2}.

To identify opportunities for outcomes improvement, registries and groups like the International Consortium for Healthcare Outcomes Measurement (ICHOM) have attempted to standardize and compare observational data across hospitals³. These comparative studies often rely on risk adjustment algorithms to account for clinical differences in patient populations⁴. In analyzing a changing medical landscape with more multi-morbidity patients, reliable morbidity data has increasingly become a focal point for fair benchmarking as part of the shift towards value-based healthcare (VBHC).

Accurate inclusion of (co)morbidities in large data sets has proven to be a vexing problem. While morbidity plays a crucial role in risk adjustment, risk stratification and outcomes measurement, there is little consensus regarding the best source of this data. Comparisons of morbidity data from different sources have displayed significant variations⁵. Notable inconsistencies have been observed when morbidity data is collected from administrative sources, such as claims data⁶. Administrative data generally under-reports comorbid conditions, leading to a lack of accounting for overall level of sickness of the patient⁷⁻⁹. While some studies have shown more accurate information in hospital chart reviews, concerns arise regarding the burden of collection and the feasibility of wide-scale use^{10,11}.

To obtain more accurate morbidity data feasibly, clinicians have increasingly turned to patient-reported instruments as a potential alternative^{12,13}. The objective of this study was to provide a comprehensive evidence base of validated patient-reported morbidity instruments to aid in the selection of these instruments for use in clinical practice. While many disease-specific morbidity instruments exist, our study examined questionnaires applicable to the broader patient population, so as to allow for broader implementation across a healthcare system.

Methods

Design & Rationale

Risk adjustment for comparison of outcomes data across (international) health care centers relies on accurate capture of predictor variables such as extent of morbidity. As standard outcome sets could be used among health institutions with no or different electronic medical record and administrative data structure, a systematic review was conducted according to the PRISMA guidelines¹⁴ of studies about the development and/or (subsequent) validation of self-reported comorbidity assessments.

Literature Search

An exhaustive search strategy was developed in Embase.com by a medical librarian experienced in systematic review searches¹⁵. To retrieve articles about the validation of questionnaires on comorbidity, the search strategy combined thesaurus terms (Emtree terms for Embase and MeSH terms for Medline) with terms in title and or abstract for three elements: comorbidity, questionnaires and validation or reliability.

The search strategy for Embase was optimized to find all potentially relevant terms and then translated to Medline (Ovid), Cochrane CENTRAL and Web of Science Core Collection¹⁶. Additional references were retrieved from Google Scholar (the first 100 references as sorted by relevance) and literature lists of relevant reviews and included references. Abstracts needed to be in English but there were no restrictions to language of the manuscript or country of publication in the search strategy. The databases were last searched on March 5th, 2018. The full search strategies for all databases is included in the online Appendix.

Study Selection

The inclusion criterion for studies was a primary focus on the development and/or (subsequent) validation (e.g. reliability and prediction/ association with outcomes) of an instrument for collecting information on the presence of morbidity directly from the patient.

The exclusion criteria were:

- Lack of any methodological description of instrument development (validation and/ or reliability of instrument);
- Description of the use of a patient-reported comorbidity questionnaire for risk-adjustment purposes or deriving health utilities; and
- Focus in the patient-reported morbidity instrument on a subset of specific conditions (based on nosological criteria), and thus not generalizable to a larger patient population (e.g. a list of mental health comorbidities for psychiatric patients).

The search results were deduplicated¹⁷ and imported thereafter in Covidence (www.covidence.org, Melbourne, Australia), a Cochrane technology web-based platform, specifically to screen and track articles through the inclusion and exclusion criteria process of a systematic review. In Covidence, the titles and abstracts of each reference were independently screened for relevance by two reviewers (AO and RB). The screening phase was conducted in the following order:

1. The first screening was based on title and abstract. In the event that the article's aim did not meet the inclusion criteria but nevertheless mentioned a patient-reported medical comorbidity questionnaire, the full text was reviewed to determine the instrument used and references were further screened for any potential missing article(s) on the comorbidity questionnaire.
2. The second screening was based on the full-text assessment of retained articles. Studies that still did not meet our inclusion criterion, were subsequently removed.

If authors AO and RB had any disagreements on article eligibility at first screening, the study in question would be screened in full-text version. Consensus on the inclusivity of selected articles was ultimately reached between both authors.

Included Comorbidities

The number of comorbidities as well as a list of included comorbid conditions was evaluated for every survey instrument. Additional survey questions, such as those evaluating the condition severity (impact on daily activities/ functional status) or medication use for a comorbidity, were also noted.

Reliability

Measures of reliability (e.g. test-retest reliability, patient-report vs. other data sources such as medical record, administrative data, or laboratory testing) at either item-level or overall instrument level were catalogued for all studies. Due to the anticipated heterogeneity in the reporting between the studies, both reliability-specific (intra-class correlation coefficient (ICC) and Kappa (κ) values) as well as other measures of the morbidity instrument's performance (Spearman's correlation coefficient, sensitivity/ specificity and positive and negative predictive values) were included. The kappa values were measured based on presence of the condition in the self-reported instrument versus the medical record (or administrative record) which was considered gold standard. The papers generally discuss that reporting a condition by self-report that is absent in the medical record could imply a deficiency with the medical record as well. Kappa values > 0.80 indicate excellent agreement, $0.61 - 0.80$ good agreement, $0.41 - 0.60$ moderate agreement, $0.21 - 0.40$ fair agreement, and < 0.20 poor agreement¹⁸. Spearman correlation coefficients are categorized in ≤ 0.20 (poor), $0.30 - 0.59$ (fair), $0.60 - 0.79$ (moderate) and > 0.80 (strong)¹⁹.

Evaluated Outcomes

All instances where patient-reported morbidity instruments were used to assess association with or predict certain outcomes as part of the validation study, were documented for this review. Examples of outcome metrics included are mortality, disease response, patient-reported outcome measures (PROMs), adverse events and other events of interest, and health care utilization/ costs as per the categories of outcomes used in the Agency for Healthcare Research and Quality Outcome Measure Framework²⁰.

Questionnaire Length, Duration, Responsiveness & Utilization

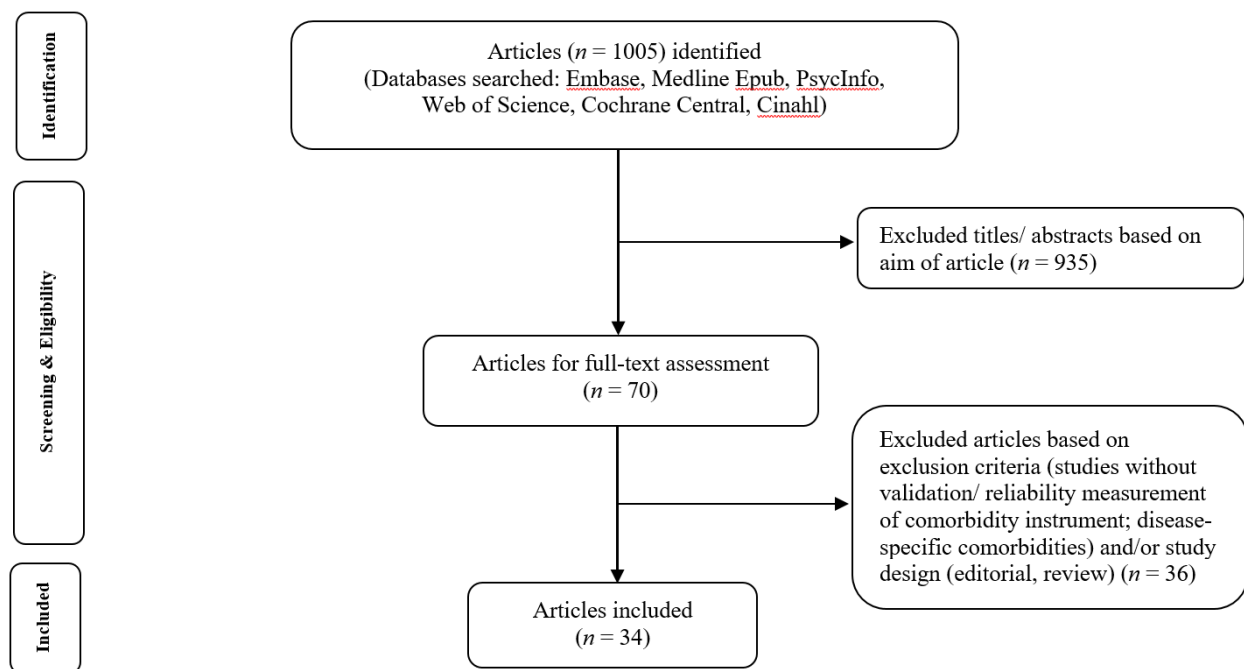
Length (number of items/ questions) of instrument and duration of completion was documented, where available, as well as route of administration (self-administration vs. administration by a clinical or research associate). Finally, the number of times the paper had been cited in Web of Science was also documented.

Results

Included Studies

Figure 1 details the search and inclusion strategy. 1005 studies met our search criteria; 70 studies met our criteria for inclusion in the full text assessment. 36 studies were eliminated after full-text review, leaving 34 studies for inclusion in this systematic review.

A summarized overview of all included articles in this systematic review are included in Table 1. Descriptive characteristics, reliability, validity and evaluated outcomes of morbidity instruments are shown in Table 2.



Flowchart of relevant article selection

Included Patient-Reported Morbidity Instruments

Ten original patient-reported morbidity instruments were identified, with most of these development studies being conducted in the United States. The instruments considered originals

were: Self-Reported Charlson Comorbidity Index (SR-CCI)¹¹, Self-Administered Comorbidity Questionnaire (SCQ)²¹, Disease Burden Morbidity Assessment (DBMA)²², Comorbidity Symptom Scale (CmSS)²³, Patient Self-Administered Health History Questionnaire²⁴, Multi-Morbidity Assessment Questionnaire for Primary Care (MAQ-PC)²⁵, Patient-Based Comorbidity Index (CI)²⁶, Health Impact Index (HII)²⁷, Seattle Index of Co-morbidity (SIC)²⁸, and an unnamed prognostic index (including comorbidities)²⁹. The SR-CCI and SCQ instruments were the most frequently cited instruments. Other included articles were either translation and cross-cultural adaptation studies, variations of these questionnaires (e.g. with a small number of items added or removed), or validation studies.

Presence of Specific Conditions & Related Assessments

Conditions that were most commonly included in the patient-reported morbidity instruments were diabetes, hypertension, myocardial infarction and stroke (Table 2). The question regarding the presence or absence of specific comorbidities was presented in multiple ways, including “Do you have or have you ever had...?”^{22,27,29-32}, “Has a doctor ever told you that you have...?”^{5,25,28,29,33-35} and “Do have you any of the following problems?”^{21,35-37}. Most instruments had close-ended response alternatives per condition listed. Some questionnaires had an additional (free-text) item for patients to report additional comorbidities that were not listed in the instrument^{21,24,38-40}.

Several instruments included additional questions regarding severity of the conditions like “Does it limit your activities?”^{5,21,22,32,36,37,40}, and/or active treatment like “Do you receive treatment for it?”^{5,21,33,36,37}.

Instrument Administration, Length, Duration & Responsiveness

Nineteen studies had self-administered questionnaires (either in mailed/ written or electronic form)^{5,21,22,24,27,28,30,32,33,36-38,40-46}, while 8 studies had questionnaires being administered verbally by a clinical/ research associate (either face-to-face or by phone)^{12,23,26,29,35,47,48} and 4 studies reported both administration methods^{9,11,31,49}. Of the associate-administered surveys, it was not clear if clarifying questions were allowed or utilized in nearly all studies. Five studies^{9,11,31,45,49} had comorbidity questionnaires that could both be self-

administered or administered by an interviewer (e.g. associate or other proxy) if needed (e.g. patient illiteracy).

Length of the instruments varied from 4 items³¹ to 195 items⁴⁴ (divided over multiple physical and mental sections). Nine studies mentioned the duration to complete the questionnaire, which ranged from 1 minute⁴¹ to 45 minutes³⁵. Response rates were provided in nine studies ranging from 28%²² to 99%³⁶.

Reliability & Concordance with Other Data Sources

Test-retest reliability was described in 7 studies^{5,11,21,23-25,40} (data not shown), mostly measured by the intra-class correlation or Spearman's correlation coefficients. The amount of patients on which it was tested ranged from 26^{11,21} – 103²⁵. The interval period between both measurements varied from 24 hours^{11,21,24} to 4 weeks²³. The overall Spearman's correlation coefficients for patient-reported comorbidity questionnaires ranged from 0.73¹¹ (moderate reliability) to 0.87²³ (strong reliability), while the intra-class correlation coefficients ranged from 0.86⁴⁰ to 0.97²⁵.

Whole instrument and item-level concordance of patient-reported morbidity scores with information from other data sources, either medical records or medical record-derived comorbidity indices, were most frequently assessed (Table 2). Spearman correlation coefficients for the relation between patient-reported morbidity scores and composite scores from other data sources ranged from $r = 0.24$ (14 conditions)³⁷ to $r = 0.70$ (18 conditions)¹¹. In studies measuring Kappa coefficients, κ values were notably higher for agreement with medical records (κ range: 0.56 – 0.69)^{24,33} as opposed to agreement with medical record-derived morbidity indices (κ range: 0.37 – 0.50)^{43,45}. Administrative data was also used as a comparative data source in a number of studies^{9,35,38,39,41}, which generally demonstrated poor agreement^{9,35,39}.

Item-level (single condition) was the most commonly reported form of concordance assessment, mostly measured against medical records or derived morbidity indices. A striking observation was that diabetes, as a comorbidity questionnaire item, had the highest Kappa value across included studies^{30,31,33-36,41,42,45,48,49}. Most included studies had a substantially wide κ value range^{21,30,33-37,41,43,45-49}, in general from 0.66²¹ to 0.86³⁷.

Association with Health & Health Care Outcomes

A number of studies assessed the association between patient-reported instrument scores and mortality, patient reported outcome measures and health care utilization. As none of the included studies evaluated their instrument against *all* three outcomes, we provided some examples in this paragraph to demonstrate the directionality of the associations.

Some studies assessed the relationship between patient-reported instrument scores with mortality and/ or survival^{26,28,29,31,41,43,44}. Habbous et al.⁴³ demonstrated a significant relation between the patient-reported Charlson Comorbidity Index and overall survival, with the presence of at least two comorbidities being associated with worse survival (HR = 1.62, p = 0.003). However, this relation was stronger for the (non-patient reported) medical record-derived CCI (HR = 2.60). Only a few studies developed prediction models for (all-cause) mortality with patient-reported comorbidity instruments, either in combination with other predictors such a demographic variables²⁹ or by themselves^{28,41}. Fan et al.²⁸ developed a prediction model with the Seattle Comorbidity Index and estimated an AUC = 0.71 for all-cause mortality at 2 years follow-up, while Lee et al.²⁹ estimated an AUC = 0.82 of a different model (including sex, age, 6 comorbidities, 4 functional measures) for all-cause mortality at 4 years follow-up.

The most commonly reported patient-reported outcome measure, evaluated with patient-reported morbidity instruments, was the 36-item Short Form Health Survey (SF-36)^{21,22,26,37,39,49}. One study reported a higher patient-reported CCI score being negatively associated with SF-36 scores⁴⁹, while another reported the number of self-reported conditions in the DBMA instrument being significantly correlated to the SF-36 similar to the medical-record derived CCI²². Selim et al.²⁶ also demonstrated a negative association between a high Patient-Based Comorbidity Index score and SF-36 scores. Due to heterogeneous reports on the association and correlation units between both measures in the included studies, there was no clear consistency observed in the (direction of the) association and/or correlation.

Several studies also analyzed the relationship between patient-reported comorbidity instruments and health care utilization outcomes (e.g. (re)hospitalizations, emergency room visits, medical costs)^{9,11,21,39,41}. Susser et al.⁹ estimated an AUC = 0.68 and AUC = 0.67 in predicting number of hospital days and emergency room visits respectively. Katz et al.¹¹ estimated weak correlations between the patient-reported comorbidity CCI and health care utilization outcomes, while Sangha et al.²¹ reported similar weak associations with the Self-Report Comorbidity Questionnaire.

Impact of Demographic Factors on Survey Validity and Reliability

Thirteen studies reported associations between certain patient characteristics and concordance of comorbidity questionnaires with other sources of comorbidity data. Most studies reported higher age being significantly associated with lower agreement between patient-reported and medical-record derived comorbidity data^{24,34,35,42,43,45-48}. However, Katz et al.¹¹, Vigen et al.³¹ and Horton et al. did not observe a significant association. In terms of reliability, Klabunde et al.⁵ found a significant association between age ≥ 65 years and inconsistent response patterns between baseline and subsequent surveys. Higher concordance between patient-reported and other comorbidity data sources was also associated with higher education levels^{11,35,42,48}. This association was also observed for higher socio-economic status⁵. In contrast, Vigen et al.³¹ only reported this association for myocardial infarction. Simpson et al.⁴⁶ reported that education level did not impact reliability with Kappa values remaining unaltered after adjustment. Some studies^{34,35,48} also reported concordance being significantly influenced by gender, while others³³ did not observe a significant observation.

Discussion

Risk adjustment for benchmarking of healthcare outcomes across multiple hospitals is dependent upon accurate reporting of case-mix factors such as patient morbidity (comorbid conditions). Clinicians have increasingly considered patient-reported comorbidity instruments as a potential alternative to the laborious review of medical and/ or administrative records, or as a method to standardize the collection of data for important conditions across hospitals. Previously published research has looked at (multi-)morbidity measures in primary care or community population settings⁵⁰ but, to our knowledge, this is the first systematic literature review focusing on the development and/ or (subsequent) validation of (general) patient-reported morbidity instruments (either indices or ad hoc lists of conditions).

Ten original patient-reported comorbidity instruments were found, as well as additional variations on these original studies. The most frequently cited instruments were the Self-Reported Charlson Comorbidity Index (SR-CCI) and the Self-Administered Comorbidity Questionnaire (SCQ), with the SR-CCI demonstrating stronger item-level reliability, overall

reliability and overall agreement, but similar correlations with health care utilization parameters. The number of items varied substantially from instrument to instrument. Most studies evaluated the accuracy of self-reported comorbid conditions for individual items versus another data source, most commonly medical records review. Agreement with the medical record, which was generally used as the gold standard, varied substantially based on the comorbid condition listed within all of the reviewed survey instruments. The kappa values regarding item-level validity were generally not used to eliminate questions with low reliability, leading to large intra-instrument variability. This variation by comorbid condition was thought by authors to be due to accurate medical record data with missed diagnoses by the patient as well as accurate reporting by the patient with missed data in the medical record. Authors postulated that disease items with low reliability included diseases that are 'resolved' (in the past)³¹, those that are controlled with treatment (e.g. hypertension)³¹, those without symptoms⁴⁶, those with complex diagnostic criteria and/or ambiguous disease categories (e.g. heart diseases such as atherosclerosis or heart failure)^{31,49}, or with confusing or overlapping names (e.g. arthritis vs. osteoarthritis vs. rheumatoid arthritis)^{33,45,46,49}. Violán et al.⁵¹ demonstrated similar results in a cross-sectional study comparing morbidity prevalence between electronic health records and health surveys: self-reported morbidity prevalence was higher among younger patients and for symptomatic conditions. Diseases with clear definitions (e.g. diabetes) and that required ongoing treatment had higher agreement with other data sources and were most accurately reported by patients^{35,43,46}. Even in those cases, there may be disagreement such as if people with pre-diabetes classify themselves as diabetic or people with non-insulin dependent diabetes consider themselves not diabetic. Agreement between patient-reported morbidity instruments and administrative data was usually poor, with the limitation generally listed in some studies that administrative data may under-report the presence of comorbid conditions ('under-coding') more than the medical record.

This systematic review highlights the lack of information on the predictive validity of comorbidity data; a subset of included studies examined the predictive capabilities of the patient-reported comorbidity questionnaires/ indices for outcomes such as mortality, patient-reported outcome measures (such as functional status or general health-related quality of life), or healthcare utilization and costs. The patterns of correlations between patient-reported comorbidity data and mortality were as hypothesized: higher patient-reported comorbidity scores

were associated with poorer overall survival⁴³. The predictive ability of patient-reported comorbidity indices were moderate to good (AUC's > 0.70) for all-cause mortality, regardless if only comorbidities were used alone or other variables were added to the model^{28,29,41}.

Unsurprisingly, higher patient-reported comorbidity scores were significantly correlated *and* associated with lower health-related quality of life scores (as measured by the SF-36)^{21,22,26,37,39,49}. Patient-reported comorbidity scores had low positive (Spearman's $r < 0.50$) correlations with many healthcare utilization measures (e.g. hospitalizations, length of hospital stay, prescribed medications)^{11,21}, and had poor discriminative ability (AUC's between 0.60 – 0.70) for these measures as well^{9,28}. As comorbidity measures can influence outcomes and interpretation while comparing treatment strategies or hospitals, it is essential that a comorbidity measure is validated for the population *and* outcome of interest.

Gold standard survey instrument development generally begins with a systematic exploration of potential topics to include in survey questions, followed by development of a large number of questions which overlap and then reduce to a smaller number of higher performing questions based on field testing^{52,53}. For example, a question on the presence or absence of myocardial infarction could be asked several ways, and the question that proves most valid and reliable in testing is retained while the other forms of the question are discarded. Although seven studies mentioned that their questionnaires were rephrased to optimize clarity and comprehension^{21-25,33,40}, there were no studies that started intentionally with a larger question bank that was then reduced down to reliable and valid questions.

This study should be interpreted in the context of its limitations. Although an extensive search in five electronic database was conducted with the help of an experienced librarian, the possibility of missing studies cannot be excluded. Comorbidity data is collected in a large proportion of medical studies, and therefore inclusion and exclusion criteria had to be defined so to allow for reasonable identification of self-reported comorbidity instruments. The goal was to identify studies that primarily focused on instrument development or validation. It is possible that comorbidity questionnaires that were used but did not have any description of their development, validity or reliability in the abstract or title could have been missed in the present search strategy, such as instruments in the grey literature. Additionally, studies that may have subsequently used information from a comorbidity instrument to predict an outcome in a general study without mention of the instrument itself in the abstract could have been missed with the

search strategy. The presence of two independent reviewers to conduct the study selection, data extraction and overall interpretation added to the accuracy of the review process.

This systematic review was inspired by international efforts (e.g. International Consortium for Health Outcomes Measurement (ICHOM)³, United States Agency for Healthcare Research and Quality (AHRQ)⁵⁴, Registry of Patient Registries project⁵⁵) to harmonize collection and format of outcome measures. These outcome measures can include suggested outcome domains, measurement tools and predictor variables for risk adjustment for a given medical condition. Among these risk-adjustment factors are morbidity variables, which can be collected and analyzed using various methods. Minimal standards for consistent capture of morbidity data are essential for fair benchmarking for VBHC or public reporting or outcomes. Patient-reported morbidity instruments can be used internationally and among hospitals with no or different electronic medical record and administrative data structure. Additionally, these instruments could be used to bolster the medical record. Future research should focus on the capture of complete morbidity data for the purposes of more robust risk adjustment, a key component of fair benchmarking for VBHC.

Conclusions

In summary, this systematic review found ten self-reported morbidity instruments, with the Patient-Reported Charlson Comorbidity Index¹¹ and the Self-Administered Comorbidity Questionnaire²¹ being the most frequently cited instruments. Within each included instrument, there was significant variability in reliability of patient-reported comorbidities based on the comorbid condition. Further research is needed to determine if patient-reported comorbidity data should be used to bolster medical records data or as a stand-alone entity for risk adjustment of observational outcomes data.

Acknowledgments

The authors would like to acknowledge Wichor M. Bramer (Biomedical Information Specialist, Medical Library, Erasmus MC) for his support in the literature search.

Authors A. Oemrawsingh and J.A. Hazelzet were supported by a grant from the Federation of Dutch University Medical Centers (NFU). R.W. Bergmark received grants from the Gliklich Healthcare Innovation Scholar Fund/ Massachusetts Eye and Ear Infirmary and from the American Board of Medical Specialties during the development of this paper.

Supplemental Material

Supplementary data associated with this article can be found in the online version at <https://doi.org/10.1016/j.jval.2020.02.006>.

References

1. Fortin M, Soubhi H, Hudon C, Bayliss EA, van den Akker M. Multimorbidity's many challenges. *BMJ : British Medical Journal*. 2007;334(7602):1016-1017.
2. Valderas JM, Starfield B, Sibbald B, Salisbury C, Roland M. Defining Comorbidity: Implications for Understanding Health and Health Services. *Annals of Family Medicine*. 2009;7(4):357-363.
3. Obbarius A, van Maasakkers L, Baer L, et al. Standardization of health outcomes assessment for depression and anxiety: recommendations from the ICHOM Depression and Anxiety Working Group. *Qual Life Res*. 2017;26(12):3211-3225.
4. Elixhauser A, Steiner C, Harris DR, Coffey RM. Comorbidity Measures for Use with Administrative Data. *Medical Care*. 1998;36(1):8-27.
5. Klabunde CN, Harlan LC, Warren JL. Data Sources for Measuring Comorbidity: A Comparison of Hospital Records and Medicare Claims for Cancer Patients. *Medical Care*. 2006;44(10):921-928.
6. Klabunde CN, Warren JL, Legler JM. Assessing Comorbidity Using Claims Data: An Overview. *Medical Care*. 2002;40(8):IV-26-IV-35.
7. de Groot V, Beckerman H, Lankhorst GJ, Bouter LM. How to measure comorbidity. a critical review of available methods. *J Clin Epidemiol*. 2003;56(3):221-229.

8. Iezzoni LI, Foley SM, Daley J, Hughes J, Fisher ES, Heeren T. Comorbidities, complications, and coding bias. Does the number of diagnosis codes matter in predicting in-hospital mortality? *Jama*. 1992;267(16):2197-2203.
9. Susser SR, McCusker J, Belzile E. Comorbidity information in older patients at an emergency visit: self-report vs. administrative data had poor agreement but similar predictive validity. *J Clin Epidemiol*. 2008;61(5):511-515.
10. Preen DB, Holman CDAJ, Lawrence DM, Baynham NJ, Semmens JB. Hospital chart review provided more accurate comorbidity information than data from a general practitioner survey or an administrative database. *Journal of Clinical Epidemiology*. 2004;57(12):1295-1304.
11. Katz JN, Chang LC, Sangha O, Fossel AH, Bates DW. Can comorbidity be measured by questionnaire rather than medical record review? *Med Care*. 1996;34(1):73-84.
12. Olomu AB, Corser WD, Stommel M, Xie Y, Holmes-Rovner M. Do self-report and medical record comorbidity data predict longitudinal functional capacity and quality of life health outcomes similarly? *BMC Health Services Research*. 2012;12(1):398.
13. Katherine F, Amy T, Nicole B, Sigrid C, Andrew JV. Comparison of Physician-Documented Versus Patient-Reported Collection of Comorbidities Among Patients With Prostate Cancer Upon First Visit to the Urology Clinic. *JCO Clinical Cancer Informatics*. 2018(2):1-10.
14. Moher D, Liberati A, Tetzlaff J, Altman DG, Group P. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med*. 2009;6(7):e1000097.
15. Bramer WM, Rethlefsen ML, Mast F, Kleijnen J. Evaluation of a new method for librarian-mediated literature searches for systematic reviews. *Res Synth Methods*. 2017.
16. Bramer WM, Rethlefsen ML, Kleijnen J, Franco OH. Optimal database combinations for literature searches in systematic reviews: a prospective exploratory study. *Syst Rev*. 2017;6(1):245.
17. Bramer WM, Giustini D, de Jonge GB, Holland L, Bekhuis T. De-duplication of database search results for systematic reviews in EndNote. *J Med Libr Assoc*. 2016;104(3):240-243.
18. Altman DG. *Practical statistics for medical research*. CRC press; 1990.

19. Akoglu H. User's guide to correlation coefficients. *Turk J Emerg Med.* 2018;18(3):91-93.
20. Gliklich RE, Leavy MB, Karl J, Campion DM, Levy D, Berliner E. A framework for creating standardized outcome measures for patient registries. *J Comp Eff Res.* 2014;3(5):473-480.
21. Sangha O, Stucki G, Liang MH, Fossel AH, Katz JN. The Self-Administered Comorbidity Questionnaire: a new method to assess comorbidity for clinical and health services research. *Arthritis Rheum.* 2003;49(2):156-163.
22. Bayliss EA, Ellis JL, Steiner JF. Subjective assessments of comorbidity correlate with quality of life health outcomes: initial validation of a comorbidity assessment instrument. *Health Qual Life Outcomes.* 2005;3:51.
23. Crabtree HL, Gray CS, Hildreth AJ, O'Connell JE, Brown J. The Comorbidity Symptom Scale: a combined disease inventory and assessment of symptom severity. *J Am Geriatr Soc.* 2000;48(12):1674-1678.
24. Boissonnault WG, Badke MB. Collecting health history information: the accuracy of a patient self-administered questionnaire in an orthopedic outpatient setting. *Phys Ther.* 2005;85(6):531-543.
25. Pati S, Hussain MA, Swain S, et al. Development and Validation of a Questionnaire to Assess Multimorbidity in Primary Care: An Indian Experience. *Biomed Res Int.* 2016;2016:6582487.
26. Selim AJ, Fincke G, Ren XS, et al. Comorbidity assessments based on patient report: results from the Veterans Health Study. *J Ambul Care Manage.* 2004;27(3):281-295.
27. Lorem GF, Schirmer H, Emaus N. Health Impact Index. Development and Validation of a Method for Classifying Comorbid Disease Measured against Self-Reported Health. *PLoS One.* 2016;11(2):e0148830.
28. Fan VS, Au D, Heagerty P, Deyo RA, McDonell MB, Fihn SD. Validation of case-mix measures derived from self-reports of diagnoses and health. *J Clin Epidemiol.* 2002;55(4):371-380.
29. Lee SJ, Lindquist K, Segal MR, Covinsky KE. Development and validation of a prognostic index for 4-year mortality in older adults. *Jama.* 2006;295(7):801-808.

30. Gad BV, Higuera CA, Klika AK, Elsharkawy KA, Barsoum WK. Validity of patient-reported comorbidities before total knee and hip arthroplasty in patients older than 65 years. *J Arthroplasty*. 2012;27(10):1750-1756 e1751.
31. Vigen C, Kwan ML, John EM, et al. Validation of self-reported comorbidity status of breast cancer patients with medical records: the California Breast Cancer Survivorship Consortium (CBCSC). *Cancer Causes Control*. 2016;27(3):391-401.
32. Wijers IGM, Ayala A, Rodriguez-Blazquez C, Rodriguez-Laso A, Rodriguez-Rodriguez V, Forjaz MJ. Disease burden morbidity assessment by self-report: Psychometric properties in older adults in Spain. *Geriatr Gerontol Int*. 2017;17(7):1102-1108.
33. Horton M, Rudick RA, Hara-Cleaver C, Marrie RA. Validation of a self-report comorbidity questionnaire for multiple sclerosis. *Neuroepidemiology*. 2010;35(2):83-90.
34. Merkin SS, Cavanaugh K, Longenecker JC, Fink NE, Levey AS, Powe NR. Agreement of self-reported comorbid conditions with medical and physician reports varied by disease among end-stage renal disease patients. *J Clin Epidemiol*. 2007;60(6):634-642.
35. Iecovich E, Biderman A. Concordance between self-reported and physician-reported chronic co-morbidity among disabled older adults. *Can J Aging*. 2013;32(3):287-297.
36. Robinski M, Strich F, Mau W, Girndt M. Validating a Patient-Reported Comorbidity Measure with Respect to Quality of Life in End-Stage Renal Disease. *PLoS One*. 2016;11(6):e0157506.
37. Stolwijk C, van Tubergen A, Ramiro S, et al. Aspects of validity of the self-administered comorbidity questionnaire in patients with ankylosing spondylitis. *Rheumatology (Oxford)*. 2014;53(6):1054-1064.
38. Lucke T, Herrera R, Wacker M, et al. Systematic Analysis of Self-Reported Comorbidities in Large Cohort Studies - A Novel Stepwise Approach by Evaluation of Medication. *PLoS One*. 2016;11(10):e0163408.
39. Voaklander DC, Kelly KD, Jones CA, Suarez-Almazor ME. Self Report Co-Morbidity and Health Related Quality of Life – A Comparison with Record Based Co-Morbidity Measures. *Social Indicators Research*. 2004;66(3):213-228.
40. Poitras ME, Fortin M, Hudon C, Haggerty J, Almirall J. Validation of the disease burden morbidity assessment by self-report in a French-speaking population. *BMC Health Serv Res*. 2012;12:35.

41. Chaudhry S, Jin L, Meltzer D. Use of a self-report-generated Charlson Comorbidity Index for predicting mortality. *Med Care*. 2005;43(6):607-615.
42. De-loyde KJ, Harrison JD, Durcinoska I, Shepherd HL, Solomon MJ, Young JM. Which information source is best? Concordance between patient report, clinician report and medical records of patient co-morbidity and adjuvant therapy health information. *J Eval Clin Pract*. 2015;21(2):339-346.
43. Habbous S, Chu KP, Harland LT, et al. Validation of a one-page patient-reported Charlson comorbidity index questionnaire for upper aerodigestive tract cancer patients. *Oral Oncol*. 2013;49(5):407-412.
44. Md Yusof MY, Horan MA, Jones M, McInnes L, Rabbitt PM, Pendleton N. Developing a self-reported comorbidity index to predict mortality of community-dwelling older adults. *Arch Gerontol Geriatr*. 2010;50(3):e63-67.
45. Mukerji SS, Duffy SA, Fowler KE, Khan M, Ronis DL, Terrell JE. Comorbidities in head and neck cancer: agreement between self-report and chart review. *Otolaryngol Head Neck Surg*. 2007;136(4):536-542.
46. Simpson CF, Boyd CM, Carlson MC, Griswold ME, Guralnik JM, Fried LP. Agreement between self-report of disease diagnoses and medical record validation in disabled older women: factors that modify agreement. *J Am Geriatr Soc*. 2004;52(1):123-127.
47. Corser W, Sikorskii A, Olomu A, Stommel M, Proden C, Holmes-Rovner M. "Concordance between comorbidity data from patient self-report interviews and medical record documentation". *BMC Health Serv Res*. 2008;8:85.
48. Hansen H, Schafer I, Schon G, et al. Agreement between self-reported and general practitioner-reported chronic conditions among multimorbid patients in primary care - results of the MultiCare Cohort Study. *BMC Fam Pract*. 2014;15:39.
49. Ng X, Low AH, Thumboo J. Comparison of the Charlson Comorbidity Index derived from self-report and medical record review in Asian patients with rheumatic diseases. *Rheumatol Int*. 2015;35(12):2005-2011.
50. Huntley AL, Johnson R, Purdy S, Valderas JM, Salisbury C. Measures of multimorbidity and morbidity burden for use in primary care and community settings: a systematic review and guide. *Ann Fam Med*. 2012;10(2):134-141.

51. Violán C, Foguet-Boreu Q, Hermosilla-Pérez E, et al. Comparison of the information provided by electronic health records data and a population health survey to estimate prevalence of selected health conditions and multimorbidity. *BMC Public Health*. 2013;13(1):251.
52. Sprangers MA, Cull A, Bjordal K, Groenvold M, Aaronson NK. The European Organization for Research and Treatment of Cancer. Approach to quality of life assessment: guidelines for developing questionnaire modules. EORTC Study Group on Quality of Life. *Qual Life Res*. 1993;2(4):287-295.
53. Farnik M, Pierzchała WA. Instrument development and evaluation for patient-related outcomes assessments. *Patient Relat Outcome Meas*. 2012;3:1-7.
54. Leroy L, Bayliss E, Domino M, et al. The Agency for Healthcare Research and Quality Multiple Chronic Conditions Research Network: overview of research contributions and future priorities. *Med Care*. 2014;52 Suppl 3:S15-22.
55. Gliklich RE DN, Leavy MB, editors. Registries for Evaluating Patient Outcomes: A User's Guide [Internet]. 3rd edition. Rockville (MD): Agency for Healthcare Research and Quality (US); 2014 Apr. 1, Patient Registries. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK208643/>.
56. McCusker J, Verdon J, Tousignant P, de Courval LP, Dendukuri N, Belzile E. Rapid emergency department intervention for older people reduces risk of functional decline: results of a multicenter randomized trial. *J Am Geriatr Soc*. 2001;49(10):1272-1281.
57. Holmes-Rovner M, Stommel M, Corser WD, et al. Does outpatient telephone coaching add to hospital quality improvement following hospitalization for acute coronary syndrome? *J Gen Intern Med*. 2008;23(9):1464-1470.
58. Spoorenberg A, van der Heijde D, de Klerk E, et al. Relative value of erythrocyte sedimentation rate and C-reactive protein in assessment of disease activity in ankylosing spondylitis. *J Rheumatol*. 1999;26(4):980-984.
59. Robinski M, Mau W, Lamprecht J, Krauth C, Girndt M. The Choice of Renal Replacement Therapy (CORETH) project: study design and methods. *Clin Kidney J*. 2014;7(6):575-581.

60. Teófilo Rodríguez J. González Cabezas A. N. Díaz Veiga P., & Rodríguez Rodríguez V. (2011). Estudio Longitudinal Envejecer en España: El proyecto ELES. Perfiles y Tendencias , 50, 1–44.
61. Kasper JD, Shapiro S, Guralnik JM, Bandeen-Roche KJ, Fried LP. Designing a community study of moderately to severely disabled older women: the Women's Health and Aging Study. *Ann Epidemiol.* 1999;9(8):498-507.
62. Young JM, Butow PN, Walsh J, et al. Multicenter randomized trial of centralized nurse-led telephone-based care coordination to improve outcomes after surgical resection for colorectal cancer: the CONNECT intervention. *J Clin Oncol.* 2013;31(28):3585-3591.
63. Schafer I, Hansen H, Schon G, et al. The German MultiCare-study: Patterns of multimorbidity in primary health care - protocol of a prospective cohort study. *BMC Health Serv Res.* 2009;9:145.
64. Potosky AL, Harlan LC, Stanford JL, et al. Prostate cancer practice patterns and quality of life: the Prostate Cancer Outcomes Study. *J Natl Cancer Inst.* 1999;91(20):1719-1724.
65. Fihn SD, McDonell MB, Diehr P, et al. Effects of sustained audit/feedback on self-reported health status of primary care patients. *Am J Med.* 2004;116(4):241-248.
66. Jacobsen BK, Eggen AE, Mathiesen EB, Wilsgaard T, Njolstad I. Cohort profile: the Tromso Study. *Int J Epidemiol.* 2012;41(4):961-967.
67. Karch A, Vogelmeier C, Welte T, et al. The German COPD cohort COSYCONET: Aims, methods and descriptive analysis of the study population at baseline. *Respir Med.* 2016;114:27-37.
68. Powe NR, Klag MJ, Sadler JH, et al. Choices for Healthy Outcomes In Caring for End Stage Renal Disease. *Seminars in Dialysis.* 1996;9(1):9-11.
69. Paleri V, Wight RG. A cross-comparison of retrospective notes extraction and combined notes extraction and patient interview in the completion of a comorbidity index (ACE-27) in a cohort of United Kingdom patients with head and neck cancer. *J Laryngol Otol.* 2002;116(11):937-941.
70. The Health and Retirement Survey. Ann Arbor: University of Michigan; 1996. Available at: http://hrsonline.isr.umich.edu/docs/sample/sho_samp.php?hfyle=ref023b&xtyp=1. Accessed January 19, 2006.

71. The Health and Retirement Survey. Ann Arbor: University of Michigan; 1999. Available at: http://hrsonline.isr.umich.edu/intro/sho_uinfo.php?hfyle=overview&xtyp=2. Accessed January 19, 2006. .
72. John EM, Phipps AI, Davis A, Koo J. Migration history, acculturation, and breast cancer risk in Hispanic women. *Cancer Epidemiol Biomarkers Prev.* 2005;14(12):2905-2913.
73. Caan B, Sternfeld B, Gunderson E, Coates A, Quesenberry C, Slattery ML. Life After Cancer Epidemiology (LACE) Study: a cohort of early stage breast cancer survivors (United States). *Cancer Causes Control.* 2005;16(5):545-556.

Table 1: Summarized Overview of Included Studies

| Patient-Reported Morbidity Instruments | Selected article and country of origin | Availability of reliability data | Evaluated outcomes | Method of questionnaire administration | Time to questionnaire completion | Number of Citations |
|---|---|---|--|---|---|----------------------------|
| Patient-Reported Charlson Comorbidity Index | Katz, 1996* ^o United States | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: + | Self- or interviewer-administered | 10 minutes | 758 |
| | Susser, 2008* Canada | Item-level: + Overall: + | Mortality: - PROM: + Healthcare utilization: - | Self-administered or filled out by proxy | - | 19 |
| | Corser, 2008* United States | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: - | Interviewer-administered | - | 53 |
| | Olomu, 2012* United States | Item-level: - Overall: - | Mortality: - PROM: + Healthcare utilization: - | Interviewer-administered | - | 23 |
| | Ng, 2015* Singapore | Item-level: + Overall: + | Mortality: - PROM: + Healthcare utilization: - | Self- or interviewer-administered | 15 minutes | 5 |
| | Habbous, 2013* Canada | Item-level: + Overall: + | Mortality: + PROM: - Healthcare utilization: - | Self-administered | - | 11 |
| | Chaudhry, 2005* United States | Item-level: + Overall: - | Mortality: + PROM: - Healthcare utilization: + | Self-administered | 1 minute | 172 |
| Self-Reported Comorbidity Questionnaire | Sangha, 2003* ^o United States | Item-level: + Overall: + | Mortality: - PROM: + Healthcare utilization: + | Self-administered | - | 757 |
| | Stolwijk, 2014* Netherlands/ Belgium | Item-level: + Overall: + | Mortality: - PROM: + Healthcare utilization: - | Self-administered | - | 17 |
| | Robinski, 2016* Germany | Item-level: + Overall: - | Mortality: - PROM: + Healthcare utilization: - | Self-administered | - | 2 |
| Disease Burden Morbidity Assessment | Bayliss, 2005 ^o United States | Item-level: + Overall: - | Mortality: - PROM: + Healthcare utilization: - | Self-administered | - | 133 |
| | Poitras, 2012 Canada | Item-level: + Overall: - | Mortality: - PROM: - | Self-administered | <15 minutes | 19 |

| | | | | | | |
|-----------------------------------|---------------------------------------|-----------------------------|--|--------------------------|----------------------|-----|
| | | | Healthcare utilization: - | | | |
| | Wijers, 2017 Spain | Item-level: - Overall: - | Mortality: - PROM: + Healthcare utilization: - | Self-administered | - | 2 |
| | Simpson, 2004 United States | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 214 |
| Comorbidity Symptom Scale | Crabtree, 2000* England | Item-level: - Overall: - | Mortality: - PROM: + Healthcare utilization: - | Interviewer-administered | <10 minutes | 29 |
| | De-loyde, 2015 Australia | Item-level: - Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 10 |
| | Gad, 2012 United States | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 3 |
| | Hansen, 2014 Germany | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | Interviewer-administered | - | - |
| | Horton, 2010 Canada/ United States | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | 11 minutes (mean) | 63 |
| Questionnaire from CALAS study | Iecovich, 2013 Israel | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: - | Interviewer-administered | 30 – 45 minutes | 1 |
| | Klabunde, 2006 United States | Item-level: - Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 114 |
| | Boissonnault, 2005 United States | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 15 |
| Seattle Index of Co- morbidity | Fan, 2002* United States | Item-level: - Overall: - | Mortality: + PROM: + Healthcare utilization: + | Self-administered | - | 123 |
| Health Impact Index | Lozem, 2016* Norway | Item-level: - Overall: + | Mortality: - PROM: + Healthcare utilization: - | Self-administered | - | 4 |
| | Lucke, 2016 Germany | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: - | Self-administered | - | 12 |

| | | | | | | |
|---|-----------------------------------|-----------------------------|--|--|--------------------|-----|
| Cornell Medical Index | Md Yusof, 2010* United Kingdom | Item-level: - Overall: + | Mortality: + PROM: - Healthcare utilization: - | Self-administered | - | 2 |
| Questionnaire from CHOICE study | Merkin, 2007 United States | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | - | - | 70 |
| Questionnaire from AHEAD study | Mukerji, 2007 United States | Item-level: + Overall: + | Mortality: - PROM: - Healthcare utilization: - | Self-administered or interviewer administered | - | 41 |
| | Paleri, 2002* United Kingdom | Item-level: + Overall: - | Mortality: - PROM: - Healthcare utilization: - | Self-administered | 8.3 minutes | 22 |
| Multi-Morbidity Assessment Questionnaire for Primary Care | Pati, 2016 India | Item-level: + Overall: - | Mortality: PROM: Healthcare utilization: | - | 20 – 25 minutes | 5 |
| | Lee, 2006* United States | Item-level: - Overall: - | Mortality: + PROM: - Healthcare utilization: - | Interviewer-administered | - | 416 |
| Self-Report Comorbidity | Voaklander, 2004* Canada | Item-level: - Overall: + | Mortality: - PROM: + Healthcare utilization: + | - | - | 12 |
| (Patient-Based) Comorbidity Index | Selim, 2004* United States | Item-level: - Overall: - | Mortality: + PROM: + Healthcare utilization: + | Interviewer-administered | - | 150 |
| Questionnaire from LACE study | Vigen, 2016 United States | Item-level: + Overall: - | Mortality: + PROM: - Healthcare utilization: - | Self- or interviewer- administered | - | 7 |

+ = described in study; - = unknown/ not described in study; * = instrument associated with an index; ° = main developmental study

Table 2: Detailed Overview of Included Studies

| | Items | Study Population (N) | Item-Level Reliability vs. Other Data Sources | Overall Instrument Reliability vs. Other Data Sources | Evaluated Outcome(s) |
|---------------------|---|--|--|--|---|
| Katz ¹¹ | <p>AIDS</p> <p>Any tumor</p> <p>Cerebrovascular disease</p> <p>Chronic pulmonary disease</p> <p>Congestive heart failure</p> <p>Connective tissue disease</p> <p>Dementia</p> <p>Diabetes (end-organ damage)</p> <p>Diabetes (mild to moderate)</p> <p>Hemiplegia</p> <p>Leukemia</p> <p>Liver disease</p> <p>Lymphoma</p> <p>Metastatic tumor</p> <p>Moderate/ Severe renal disease</p> <p>Myocardial infarction</p> <p>Peripheral vascular disease</p> <p>Ulcer disease</p> | <p>170 inpatients from 6 care units (3 medical and 3 surgical) at 1 hospital.</p> <p>Characteristics:</p> <p>Female 55%</p> <p>Mean age 65.3 years (\pm SD 8.8)</p> <p>Caucasian 82%</p> <p>College level or higher 50%</p> <p>Surgical 54%</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results:</p> <p>Kappa: 0.35 (ulcer disease/diabetes with end-organ damage) – 0.85 (leukemia)</p> <p>Sensitivity: -</p> <p>Specificity: -</p> <p>PPV: -</p> <p>NPV: -</p> <p>Agreement between self-reported CCI and medical record-derived CCI ranged from 83% (any tumor) to 100% (AIDS).</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results:</p> <p>Self-reported CCI score was higher versus medical-record derived CCI ($1.99 \pm$ SD 2.13 vs. $1.59 \pm$ SD 1.80, $p < 0.01$)</p> <p>Spearman's r range 0.63 ($p = 0.0001$) (full index) – 0.70 ($p = 0.0001$) (when the solid tumor item was excluded from the analysis)</p> | <p>Measured in 49 and 56 patients on medical and surgical service respectively.</p> <p>Results:</p> <p>Mortality: -</p> <p>PROM: -</p> <p>Health care utilization:</p> <p><i>Hospitalizations in last year:</i> Spearman's r range 0.17 – 0.31, $p < 0.05$</p> <p><i>Number of prescription medication:</i> Spearman's r range 0.26 – 0.44, $p < 0.05$</p> <p><i>Hospital charges during admission:</i> Spearman's r range 0.09 – 0.26, $p < 0.05$</p> <p><i>Length of stay:</i> Spearman's r range 0.15 – 0.20</p> |
| Susser ⁹ | As per Katz | <p>520 elderly patients ready to be discharged home from the ER. Data from a previously published RCT⁵⁶.</p> <p>Characteristics:</p> <p>Female 60%</p> <p>Age group >75 years 57%</p> | <p>Comparison: Administrative data-derived CCI</p> <p>Results:</p> <p>Kappa: Highest $\kappa = 0.55$ (chronic pulmonary disease). Individual Kappa values with range were not described for all conditions.</p> <p>Sensitivity: -</p> <p>Specificity: -</p> <p>PPV: -</p> <p>NPV: -</p> <p>Four conditions were reported more frequently by self-report (myocardial infarction, ulcer disease, diabetes with end-</p> | <p>Comparison: Administrative data-derived CCI</p> <p>Results:</p> <p>Poor agreement between self-reported and administrative data-derived CCI, indicated by an (overall) ICC=0.43 (95% CI 0.40 – 0.47).</p> | <p>Comparison: Administrative data-derived CCI</p> <p>Results:</p> <p>Mortality: -</p> <p>PROM:</p> <p><i>ADL (functional decline):</i> predictive ability of self-reported vs. administrative data-derived CCI was measured with unweighted (for sampling) AUC=0.51 vs. AUC=0.54 and weighted AUC=0.54 vs. AUC=0.50, $p > 0.05$)</p> <p>Health care utilization:</p> <p><i>Hospital days:</i> self-reported vs. administrative data-derived CCI was measured with unweighted AUC=0.63 vs. AUC=0.63 and weighted AUC=0.68 vs. AUC=0.69, $p > 0.05$)</p> <p><i>ER visits:</i> unweighted AUC=0.64 vs. AUC=0.65 and weighted AUC=0.67 vs. AUC=0.63, $p > 0.05$)</p> |

| | | | | | |
|--------------------------|-------------|--|--|--|---|
| | | | organ damage, connective tissue disease), while five (hemiplegia, mild-moderate diabetes, solid tumor, lymphoma, dementia) were more frequently reported in administrative data. | | |
| Cors er ⁴⁷ | As per Katz | <p>525 patients admitted for acute coronary syndrome in 5 hospitals.</p> <p>Characteristics: Female 36.4% Mean age 59.73 years (\pm SD 12) Caucasian 84.4% College level or higher 43.8%</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Kappa: 0.07 (CTD, RA) – 0.80 (diabetes). Only conditions with a prevalence of at least 3% (in each data source) were included in the Kappa analysis. Sensitivity: - Specificity: - PPV: - NPV: -</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Self-reported CCI (composite) scores were higher than medical record-derived CCI scores (mean $1.78 \pm$ SD 1.99 vs. mean $1.27 \pm$ SD 1.43). Correlation between self-reported and medical record-derived CCI composite scores were fair (Spearman's $r=0.57$, $p<0.01$).</p> | - |
| Olo mu ¹² | As per Katz | <p>525 patients admitted for acute coronary syndrome in 5 hospitals. Data from a previously published RCT⁵⁷.</p> <p>Characteristics: Female 36.4% Caucasian 84.4% College level or higher 43.8%</p> | - | - | <p>Comparison: Medical record-derived CCI.</p> <p>Results: Mortality: - PROM: <i>ASI (functional capacity)</i>: Prediction at 3 months was slightly better with SCQ vs. the CCI ($R^2=0.340$, $p<0.0005$ vs. $R^2=0.331$, $p<0.0035$), while it was slightly better with CCI vs. SCQ ($R^2=0.370$, $p<0.0005$ vs. $R^2=0.358$, $p<0.0005$) at 8 months. <i>EQ-5D (health-related quality of life)</i>: Only the SCQ significantly predicted EQ-5D scores at 3 and 8 months ($R^2=0.288$ and $R^2=0.265$, $p<0.0005$), whereas the CCI did not ($R^2=0.262$, $p>0.201$ and $R^2=0.245$, $p>0.132$). Health care utilization: -</p> |

| | | | | | |
|-------------------|---|--|--|---|--|
| Ng ⁴⁹ | As per Katz | <p>301 rheumatic patients from 1 tertiary hospital.</p> <p>Characteristics: Female 61.5% Median age 51 years (21-79) Chinese 68.8% College level or higher 54.7%</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Kappa: 0.189 (diabetes with end-organ damage) – 0.764 (diabetes). Kappa values was only calculated for 8/18 conditions that did not have any cell values of zero. Sensitivity: 33.3 (diabetes with end-organ damage) – 100% (myocardial infarction) Specificity: 58.9 (CTD, RA) – 99.1% (CVA) PPV: - NPV: -</p> <p>Agreement between self-reported CCI and medical record-derived CCI ranged from 74.1% (CTD/RA) to 100% (leukemia, lymphoma, metastatic solid tumor, AIDS).</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Median self-reported composite CCI scores were higher than the medical record-derived CCI scores, indicating that conditions were generally reported more frequently by self-report than EHR review. Self-reported composite CCI scores had moderate agreement (ICC=0.513, p<0.001) and strong correlation (Spearman's r=0.570, p<0.001) with the medical record-derived CCI scores.</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Mortality: - PROM: <i>SF-36 (health-related quality of life)</i>: Self-reported CCI was negatively associated with PCS (β=-2.56, p<0.001) and MCS (β=-1.24, p=0.044). Medical record-derived CCI scores had a similar trend but coefficients didn't reach statistical significance. Health care utilization: -</p> |
| Hab bous 43 | <p><i>Exposures</i> Smoking and alcohol</p> <p><i>Conditions/Diseases</i> Chronic cough/bronchitis Dementia (e.g. Alzheimer's) Diabetes (eye/kidney problems) (Past) Dialysis requirement Emphysema Heart failure Hemiplegia Hepatitis HIV/AIDS Liver disease Myocardial infarction Other joint/bone problems</p> | <p>882 head-and-neck cancer patients.</p> <p>Characteristics: Female 23% Median age 61.5 years (61-62.5) Caucasian 84%</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Kappa: 0.16 (hemiplegia) – 0.93 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: -</p> <p>Positive agreement between self-reported CCI and medical record-derived CCI ranged from 17 (hemiplegia) to 94% (diabetes). Negative</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Patient-reported CCI scores were higher than the medical record-derived CCI scores (mean 1.01 (95% CI 0.9 – 1.1) vs. 0.74 (95% CI 0.7 – 0.8), p<0.0001). Comorbidities were reported more often by patients in comparison to medical records review. Overall agreement between patient-reported CCI and medical record-derived CCI was measured as κ=0.37, which improved if CTD (κ=0.52) or</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Mortality: <i>Overall survival</i>: Both patient-reported CCI (HR 1.62 (95% CI 1.18 – 2.24), p=0.003) and medical record-derived CCI (HR 1.97 (95% CI 1.38 – 2.80), p=0.0002) were significantly associated with overall survival after multivariate (age, sex, marital status, stage and disease site) adjustment, when at least 2 comorbidities were present. PROM: - Health care utilization: -</p> |

| | | | | | |
|-------------------------------|--|--|---|---|---|
| | <p>Past cancer history</p> <p>Peripheral vascular disease</p> <p>Rheumatoid arthritis</p> <p>Serious kidney problems</p> <p>Stomach ulcers (test-proven)</p> <p>Stroke/ Mini-stroke</p> | | <p>agreement between self-reported CCI and medical record-derived CCI ranged from 84 (CTD) to 100% (dementia).</p> | <p>COPD ($\kappa=0.43$) was removed from the patient-reported CCI score.</p> | |
| <p>Chau dhry⁴¹</p> | <p>Asthma/ Emphysema/ Chronic bronchitis</p> <p>Arthritis or rheumatism</p> <p>Cancer (diagnosed within past 3 years)</p> <p>Diabetes</p> <p>Digestive problems (i.e. ulcer/colitis/ gallbladder disease)</p> <p>Heart trouble (i.e. angina/CHF/ CAD)</p> <p>HIV or AIDS</p> <p>Kidney disease</p> <p>Liver problems (cirrhosis)</p> <p>Stroke</p> | <p>7761 hospitalized general medicine patients at a single center.</p> <p>Characteristics:</p> <p>Female 62%</p> <p>Mean age 56-57 years</p> <p>African-American >80%</p> <p>MMSE score > 17</p> | <p>Comparison: Administrative data-derived CCI</p> <p>Results:</p> <p>Item-level data vs. one-year look-back:</p> <p>Kappa: 0.04 (stomach ulcer) – 0.83 (diabetes)</p> <p>Sensitivity: 44 (cancer) – 86% (diabetes, HIV/AIDS)</p> <p>Specificity: 48 (arthritis or rheumatism) – 98% (HIV/AIDS)</p> <p>PPV: 3 (stomach ulcers) – 90% (diabetes)</p> <p>NPV: 91 (asthma, emphysema or bronchitis) – 100% (HIV/ AIDS)</p> <p>Item-level data vs. index hospitalization:</p> <p>Kappa: 0.06 (arthritis or rheumatism) – 0.82 (diabetes)</p> <p>Sensitivity: 43 (cancer) – 91% (diabetes)</p> <p>Specificity: 47 (arthritis or rheumatism) – 95% (liver disease, cancer)</p> <p>PPV: 9 (arthritis or rheumatism) – 84% (diabetes)</p> <p>NPV: 93 (asthma, emphysema or bronchitis) – 99% (heart disease, kidney disease, liver disease, cancer)</p> | - | <p>The predictive power of the <i>self-reported</i> CCI was constructed with 4 different logistic regression models performed in a validation cohort (N=3870).</p> <p>Model 1: age, sex + original CCI weight</p> <p>Model 2: age, sex + study-specific CCI weight</p> <p>Model 3: age, sex, diagnosis-related group weight + original CCI weight,</p> <p>Model 4: age, sex, diagnosis-related group weight + study-specific CCI weight</p> <p>Results:</p> <p>Mortality:</p> <p><i>One-year mortality</i>: AUC's for the self-reported CCI were 0.70 (0.68 – 0.73) for model 1, 0.72 (0.70 – 0.75) for model 2, 0.75 (0.72 – 0.77) for model 3, and 0.76 (0.73 – 0.78) for model 4. AUC's were slightly less compared to the administrative data-derived CCI indices ($p<0.001$).</p> <p>PROM: -</p> <p>Health care utilization:</p> <p><i>Log total costs</i>: R^2 values for the different regression models ranged from 0.02 (models 1 & 2) – 0.33 (models 3 & 4)</p> <p><i>Log length of stay</i>: R^2 values for the different regression models ranged from 0.01 (model 1) – 0.22 (models 3 & 4)</p> |

| | | | | | |
|----------------------|---|--|---|---|---|
| | | | No statistically significant differences in Kappa values, sensitivities, specificities, or positive or negative predictive values was observed for 1-year look-back periods or index hospitalization. | | |
| Sangha ²¹ | <p>Anemia or other blood disease</p> <p>Back pain</p> <p>Cancer</p> <p>Depression</p> <p>Diabetes</p> <p>Heart disease</p> <p>High blood pressure</p> <p>Kidney disease</p> <p>Liver disease</p> <p>Lung disease</p> <p>Osteoarthritis/ Degenerative arthritis</p> <p>Other medical problems (optional)</p> <p>Rheumatoid arthritis</p> <p>Ulcer or stomach disease</p> | <p>170 hospitalized patients from 6 care units at one hospital.</p> <p>Characteristics: Female 55% Mean age 65.3 years (\pm SD 8.8) Caucasian 82% College level or higher 50%</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Kappa: 0.27 (lung disease) – 0.93 (liver disease) Sensitivity: - Specificity: - PPV: - NPV: -</p> <p>Overall agreement between the SCQ and the medical record-derived CCI ranged from 78% (heart disease) to 99% (liver disease).</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: SCQ scores were higher than the medical record-derived CCI scores (mean $5.61 \pm$ SD 4.1 vs. $1.59 \pm$ SD 2.13) SCQ had a fair correlation (Spearman's $r=0.32$) with the medical record-derived CCI, which slightly increased (Spearman's $r=0.55$) when questionnaires were truncated to only comparable items.</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results for medical patients: Mortality: - PROM: <i>SF-36 (health-related quality of life)</i>: SCQ had poor to modest correlations with SF-36 (subscale) scores at one-year follow-up, ranging from 'MCS' (Spearman's $r=-0.03$, $p>0.05$) to 'General health' (Spearman's $r=-0.39$, $p<0.0001$). Total SCQ scores explained substantial variation for most SF-36 subscales, with R^2 values ranging from 0.10 ('Social function') to 0.25 ('Physical function') in multivariate (including age, sex, ethnicity, education level, and insurance status) linear regression models. Health care utilization: <i>Hospitalizations in previous year</i>: SCQ scores correlated fairly with hospitalizations in the previous 12 months (Spearman's $r=0.21$, $p<0.01$ for medical patients; Spearman's $r=0.37$, $p<0.01$ for surgical patients) <i>Prescription medication</i>: SCQ scores also correlated moderately with number of prescriptions (Spearman's $r=0.40$ for medical patients; $r=0.55$ for surgical patients) <i>Total hospital charges</i>: SCQ scores correlated poorly with total inpatient charges (Spearman's $r=0.09$ for medical patients; $r=0.10$ for surgical patients) <i>Length of stay</i>: SCQ scores also correlated poorly with hospital length of stay (Spearman's</p> |

| | | | | | |
|------------------------|--|--|--|---|--|
| | | | | | r=0.03 for medical patients; r=0.14 for surgical patients) |
| Stolwijk ³⁷ | <p>Dutch modified version (mSCQ) of the SCQ instrument²¹</p> <p>Anemia or other blood disease Back pain Cancer Depression Diabetes Heart disease High blood pressure Kidney disease Liver disease Lung disease Osteoarthritis Other non-specified medical problems (optional; max. 3) Rheumatoid arthritis Ulcer or stomach disease</p> | <p>98 outpatients with ankylosing spondylitis. Data from the OASIS study⁵⁸.</p> <p>Characteristics: Female 29.6% Mean age 53.9 years (\pm SD 11.4) College level or higher 15.7%</p> | <p>Comparison: Medical records</p> <p>Results: Kappa: 0.14 (osteoarthritis, ulcer disease) – 1.00 (cancer). Kappa analysis included 10 conditions. Sensitivity: - Specificity: - PPV: - NPV: -</p> | <p>Comparisons: 1. Medical record-derived CCI 2. Michaud-Wolfe index</p> <p>Results: SCQ had poor to fair correlations with the medical record-derived CCI (Spearman's $r=0.24$, $p<0.05$) and Michaud-Wolfe index (Spearman's $r=0.43$, $p<0.05$) mSCQ also had moderate correlations with CCI (Spearman's $r=0.36$, $p<0.05$) and Michaud-Wolfe index (Spearman's $r=0.57$, $p<0.05$)</p> | <p>Comparisons: 1. Medical record-derived CCI 2. Michaud-Wolfe index</p> <p>Results: Mortality: - PROM: <i>BASDAI (disease activity)</i>: SCQ correlated moderately with disease activity (Spearman's $r=0.27$, $p<0.05$), while CCI correlated poorly (Spearman's $r=0.01$). SCQ was significantly associated (OR=1.73, 95% CI 1.25 – 2.40, $p<0.01$) with low disease activity (BASDAI<4). <i>BASFI (physical function)</i>: SCQ correlated moderately (Spearman's $r=0.43$, $p<0.05$) with physical function, but was significantly associated ($\beta=0.11$, 95% CI 0.03 – 0.19, $p=0.01$) with BASFI. <i>SF-36 (health-related quality of life)</i>: SCQ correlated moderately (Spearman's $r=-0.45$, $p<0.05$) with the PCS subscale, and was significantly associated ($\beta=-0.72$, $p=0.03$) with PCS. CCI and Michaud-Wolfe indices were not significantly associated with BASFI and SF-36-PCS. Health care utilization: -</p> |
| Robinski ³⁶ | German version (SCQ-G) of the SCQ instrument ²¹ | <p>780 adult end-stage renal disease patients from 55 dialysis units. Data from the CORETH project⁵⁹.</p> <p>Characteristics: Female 32.6% Mean age 63.2 years (\pm SD 15.1)</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Kappa: 0.01 (peptic ulcer disease) – 0.84 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: -</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Total SCQ-G score was moderately correlated to the medical record-derived CCI (Spearman's $r=0.27$, $p<0.01$).</p> | <p>Comparison: Medical record-derived CCI</p> <p>Results: Mortality: - PROM: <i>SF-12 (health-related quality of life)</i>: Total SCQ-G score was moderately correlated with MSC (Spearman's $r=-0.25$, $p<0.01$) and PSC (Spearman's $r=-0.49$, $p<0.01$) subscales, while CCI correlated poorly with MSC (Spearman's $r=0.06$, $p>0.05$) and moderately with PSC (Spearman's $r=-0.36$, $p<0.01$).</p> |

| | | | | | |
|-----------------------|--|---|---|---|---|
| | | | Overall agreement between the SCQ and CCI ranged from 70 (heart disease) to 95% (kidney disease, liver disease). Positive agreement between both data sources ranged from 6% (peptic ulcer disease) to 97% (kidney disease). Negative agreement ranged from 78% (heart disease) to 97% (liver disease). | | Health care utilization: - |
| Bayliss ²² | Angina/ CAD Asthma Back pain (chronic) or sciatica Bronchitis (chronic)/ COPD Cancer (diagnosed within past 5 years) Cholesterol (elevated) Colon problem (e.g. diverticulitis/ irritable bowel) Congestive heart failure Diabetes Hard of hearing Hypertension Kidney disease Nerve condition Osteoarthritis Osteoporosis Overweight Poor circulation (e.g. peripheral vascular disease) Rheumatic disease (e.g. fibromyalgia or lupus) Rheumatoid arthritis Stomach problem (e.g. gastritis/ ulcer/ reflux) Stroke Thyroid disorder Vision problem | 156 patients (≥ 65 years) from the HMO Characteristics: Female 49.4% Mean age 75 years (67-94) Caucasian 91% College level or higher 59.6% | Comparison: Medical records Results: Kappa: - Sensitivity: 35 (kidney disease) – 100% (asthma) Specificity: 61 (hard of hearing) – 100% (kidney disease, cancer) PPV: - NPV: - | Comparison: Medical records Results: Sensitivity by respondent analysis: 14 – 100% (median 75%) Specificity by respondent analysis: 59 – 100% (median 91%) | Comparisons: 1. Medical records 2. Medical record-derived CCI 3. Rx-risk score (comorbidity measure including age, gender, health insurance benefit status and a category based on diagnoses from administrative pharmacy data) Results: Mortality: - PROM: <i>SF-36 (health-related quality of life)</i> : Self-reported number of conditions (Spearman's $r=0.477$, $p<0.001$) had a similar correlation compared to the medical-record CCI ($r=0.48$, $p<0.001$) but higher compared to the RxRisk score (0.17 , $p=0.037$). <i>SF-36 (physical functioning)</i> : Self-reported conditions ($r=-0.482$, $p<0.001$) had a stronger correlation vs. CCI ($r=-0.41$, $p<0.001$) and the RxRisk score ($r=-0.18$, $p=0.035$). <i>BRFSS (depression screening)</i> : Self-reported conditions ($r=-0.24$, $p=0.003$) had a stronger correlation compared to CCI ($r=-0.12$, $p=0.14$) and the a RxRisk score ($r=-0.05$, $p=0.559$). <i>GSE (self-efficacy)</i> : Self-reported conditions ($r=-0.305$, $p<0.001$) had a stronger correlation vs. CCI ($r=-0.14$, $p=0.096$) and the RxRisk score ($r=0.10$, $p=0.234$). |

| | | | | | |
|---------------------------|---|---|--|--|--|
| | | | | | Health care utilization: - |
| Poitr as ⁴⁰ | French modified version (DBMA-Fv) of the DBMA instrument ²² : 21 of the 23 original conditions were chosen Items “Kidney disease” and “Nerve condition” were excluded in this version Item “Depression/ anxiety” was added to this version | 78 patients from 1 health center. Characteristics: Female 68% Mean age 47.4 years (\pm SD 15.9) College level or higher 70.5% | Comparison: Medical records Results: Kappa: - Sensitivity: 62.5 (angina/ CAD) – 90% (diabetes) - Specificity: 77.6 (overweight) – 98.6% (diabetes) PPV: 44.4 (overweight) – 92.9% (hypercholesterolemia) NPV: 88.7 (osteoarthritis) – 95.9% (asthma/ diabetes) | Comparison: Cumulative Illness Rating Scale Results: DBMA-Fv correlated moderately with the CIRS at baseline ($r=0.46$, 95% CI 0.26 – 0.62, $p<0.01$) and at two weeks follow-up (Spearman’s $r=0.56$, 95% CI 0.38 – 0.70, $p<0.01$). Comparison: Medical records Results: Mean sensitivity of patient-reported conditions vs. medical record review at two weeks was 73.9% (\pm SD 8.4), while mean specificity was 92.2% (\pm SD 6.7). | - |
| Wije rs ³² | Spanish modified version of the DBMA instrument ²² : 21 out of the 23 original conditions were chosen Item “Liver disease” was excluded from this version due to low prevalence Items “UTI”, “anxiety” and “memory-related disorders” were added to this version due to high due to high prevalence in older adults | 707 community-dwelling adults (≥ 65 years). Data from the ELES-PS study ⁶⁰ . Characteristics: Female 57% Mean age 74.2 years (\pm 6.6) College level or higher 17.3% | - | - | Comparison: Self-reported conditions Results: Mortality: - PROM: <i>PWI (physical functioning & perceived health status)</i> : DBMA significantly correlated stronger to physical functioning than self-reported number of conditions (Spearman’s $r=-0.56$ vs. $r=-0.51$, $p=0.0035$) <i>PWI (quality of life)</i> : DBMA significantly correlated stronger to PWI in comparison to self-reported number of conditions (Spearman’s $r=-0.41$ vs. $r=-0.35$, $p=0.0006$) <i>CES-D (depression screening)</i> : DBMA significantly correlated stronger to CES-D compared to self-reported number of conditions (Spearman’s $r=0.41$ vs. $r=0.35$, $p=0.0043$) Health care utilization: - |

| | | | | | |
|----------------------------|--|---|---|---|---|
| Sim pson ⁴⁶ | <p>Angina pectoris</p> <p>Arthritis (OA/ RA)</p> <p>Cancer</p> <p>Congestive heart failure</p> <p>Diabetes mellitus</p> <p>Disc disease</p> <p>Hip fracture</p> <p>Lung disease</p> <p>Myocardial infarction</p> <p>Osteoporosis</p> <p>Parkinson's disease</p> <p>Peripheral arterial disease</p> <p>Spinal stenosis</p> <p>Stroke</p> | <p>1002 disabled women, aged ≥ 65 years, with MMSE ≥ 18. Data from the Women's Health and Aging Study I⁶¹.</p> <p>Characteristics: Age group 65-74 years 44.2% Caucasian 71.1%</p> | <p>Comparison: Disease-specific standardized algorithms (medical history, physical examination, medication review, medical record review, X-rays, physician questionnaire)</p> <p>Results: Kappa: 0.24 (peripheral arterial disease) – 0.96 (hip fracture) Sensitivity: 22 (spinal stenosis) – 98% (stroke) Specificity: 45 (arthritis) – 100% (hip fracture, Parkinson's disease, disc disease, spinal stenosis) PPV: 0.20 (peripheral arterial disease) – 1.0 (Parkinson's disease) NPV: 0.38 (arthritis) – 1.0 (hip fracture, Parkinson's disease, cancer, stroke)</p> | - | - |
| Crab tree ²³ | <p>Angina</p> <p>Anxiety and depression</p> <p>Any other condition</p> <p>Arthritis/ Osteoporosis</p> <p>Breathlessness secondary to cardiovascular cause</p> <p>Breathlessness/ Wheeze (secondary to respiratory cause)</p> <p>Cerebrovascular disease</p> <p>Constipation</p> <p>Cough/ Sputum (secondary to COPD/ Asthma)</p> <p>Diabetes</p> <p>Diarrhea</p> <p>Epilepsy</p> <p>Hearing problems</p> <p>Pain</p> | <p>183 patients ≥ 65 years with confirmed age-related cataract (N=161) or from a geriatric day hospital (N=22)</p> | - | - | <p>Comparison: -</p> <p>Results: Mortality: - PROM: <i>NEADL (activities of daily living)</i>: CmSS correlated moderately to the NEADL (Spearman's $r=0.56$, $p<0.01$). <i>GHQ-28 (perceived health status)</i>: CmSS correlated poorly to the GHQ-28 (Spearman's $r=0.48$, $p<0.01$) <i>HAD (anxiety and depression)</i>: CmSS correlated moderately to the HAD (Spearman's $r=0.52$, $p<0.01$). Health care utilization: -</p> |

| | | | | | |
|----------------------------|---|---|---|---|---|
| | Parkinson's disease Peripheral vascular disease Side effects from medications Skin disease Unsteadiness, falls and syncope Upper gastrointestinal symptoms Urinary problems Visual problems Walking and mobility | | | | |
| De-loyd e ⁴² | Another cancer Chronic respiratory disease Depression Diabetes Heart disease Hypertension Kidney disease | 756 patients with colorectal cancer from multiple hospitals. Data from the CONNECT ⁶² RCT. Characteristics: Female 56% Age group <70 years 54% College level or higher 24% | Comparison: Clinician report Results: Kappa: 0.22 (another cancer) – 0.58 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: - Comparison: Medical records Results: Kappa: 0.34 (kidney disease) – 0.77 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: - | - | - |
| Gad ³ 0 | Amputation(s) Anemia or other blood problem(s) Asthma/ Other lung disease Back pain Blood clots or phlebitis Bowel problems Cancer Chronic skin condition Congestive heart failure Depression or anxiety Diabetes Excessive weight | 382 preoperative orthopedic patients (aged ≥ 65 years) before undergoing total knee or hip arthroplasty Characteristics: Female 65% Mean age 74 years (± SD 6.1) | Comparison: Medical records Results: Kappa: 0.00 (osteoarthritis) – 0.76 (diabetes) Sensitivity: 9 (peripheral vascular disease) – 71% (hypertension) Specificity: 44 (osteoarthritis) – 99% (diabetes) PPV: - NPV: - | - | - |

| | | | | | |
|----------------------|--|---|--|---|---|
| | Hearing loss Heart attack High blood pressure High cholesterol Kidney or urinary problems Liver/ Gallbladder disease Lupus/ Other autoimmune disease Neuromuscular disease Osteoarthritis/ Degenerative arthritis Osteoporosis Paralysis Peripheral vascular disease Previous fracture(s) Recent unwanted weight loss Rheumatoid arthritis Sleep problems Stroke Thyroid problems Ulcer/ Stomach problems Visual problems | | | | |
| Hansen ⁴⁸ | <i>For analysis (32/46 diagnosis groups):</i> Anemia Asthma/ COPD Atherosclerosis/ PAOD Cancers Cardiac arrhythmias Cardiac insufficiency Cardiac valve disorders Cerebral ischemia/ Chronic stroke Chronic cholecystitis/ Gallstones Chronic ischemic heart disease Chronic low back pain Diabetes mellitus Dizziness Gynecological problems Hemorrhoids Hypertension Hyperuricemia/ Gout Intestinal diverticulosis | 3189 multi-morbid primary care patients. Data from the Multi-Care Cohort Study ⁶³ . Characteristics: Female 59.3% Mean age 74.4 years (\pm SD 5.2) College level or higher 10.9% | Comparison: Clinician report Results: Kappa: 0.05 (gynecological problems) – 0.80 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: - | - | - |

| | | | | | |
|----------------------|--|--|--|---|---|
| | Joint arthrosis Lipid metabolism disorders Lower limb varicosis Migraine/ Chronic headache Neuropathies Osteoporosis Parkinson's disease Prostatic hyperplasia Psoriasis Renal insufficiency Rheumatoid arthritis/ Chronic polyarthritis Severe vision reduction Thyroid dysfunction Urinary tract calculi | | | | |
| Horton ³³ | Anemia Anxiety Arthritis Bipolar disorder Breast cancer Cataracts Colon cancer Depression Diabetes Epilepsy Fibromyalgia Glaucoma Heart disease Hip replacement Hyperlipidemia Hypertension Inflammatory bowel disease Irritable bowel syndrome Kidney disease Knee replacement Liver disease Lung cancer Lung disease Migraine Osteoporosis Peptic ulcer disease | 404 patients with multiple sclerosis from 2 centers. Characteristics: Female 76% Mean age 46.5 years (\pm SD 11.8) Caucasian 92% College level or higher 63.4% Relapsing-remitting MS 70.8% | Comparison: Medical records Results: Kappa: 0.19 (anemia) – 0.88 (diabetes) Sensitivity: 14 (kidney disease) – 100% (bipolar disorder, breast cancer, glaucoma, lung cancer, rheumatoid arthritis, schizophrenia, cataracts) Specificity: 87 (depression) – 100% (breast cancer, lung cancer) PPV: 0.07 (skin cancer) – 1.00 (breast cancer/ lung cancer) NPV: 0.84 (depression) – 1.00 (bipolar disorder, breast cancer, cataracts, glaucoma, lung cancer, rheumatoid arthritis, schizophrenia) | Comparison: Medical records Results: Agreement between self-reports and medical records was $\kappa=0.56$ (95% CI 0.48 – 0.64) for the presence of any physical comorbidity, and $\kappa=0.57$ (95% CI 0.48 – 0.65) for mental comorbidities. For this analysis, the questionnaire was divided into physical vs. mental comorbid conditions, and thereafter dichotomized in 0 vs. >0 comorbidities. | - |

| | | | | | |
|------------------------|---|--|---|---|---|
| | Peripheral vascular disease Rectal cancer Rheumatoid arthritis Schizophrenia Sjogren's syndrome Skin cancer Systemic lupus erythematosus Thyroid Uveitis Vitamin-B12 deficiency | | | | |
| Iecovich ³⁵ | Arthritis Cancer Cardiovascular accident Circulatory disease Diabetes Gastrointestinal disease Hypertension Myocardial infarction Osteoporosis Other heart diseases Renal problems Respiratory disease Thyroid disease | 402 disabled older patients who used adult daycare centers. Characteristics: Female 74.8% Mean age 78 years (\pm SD 7.02) Asian/ African 62.6% College level or higher 10.1% | Comparison: Medical records (including diagnostic ICD-9 codes) Results: Kappa: 0.09 (circulatory disease) – 0.76 (diabetes) Sensitivity: 22.5 (cancer) - 79.1% (diabetes) - Specificity: 73.5 (renal) – 98% (cancer) PPV: 0.36 (circulatory disease) – 0.92 (hypertension) NPV: 0.42 (hypertension) – 0.87 (thyroid disorder) | Comparison: Medical records (including diagnostic ICD-9 codes) Results: Self-reports correlated fairly with the EHR ($r=0.45$, $p<0.001$). | - |
| Klabunde ⁵ | Angina Arthritis or rheumatism Chronic Lung Disease/ Bronchitis/ Emphysema Cirrhosis/ Liver disease Congestive heart failure Depression or anxiety Diabetes Hypertension IBD/ Colitis/ Crohn's disease Myocardial infarction Stroke/ Brain hemorrhage Stomach ulcers with bleeding | 3095 prostate cancer survivors. Data from the PCOS study ⁶⁴ . Characteristics: Age group >65 years 64% Caucasian 78% College level or higher 60% | - | - | - |

| | | | | | |
|----------------------------|--|---|---|--|---|
| Boissonnault ²⁴ | <p>Anemia</p> <p>Ankylosing spondylitis</p> <p>Arterial blockage of legs</p> <p>Asthma</p> <p>Cancer</p> <p>Chemical dependency</p> <p>Deep venous thrombosis</p> <p>Degenerative osteoarthritis or wear-and-tear arthritis</p> <p>Depression</p> <p>Diabetes (diagnosed after age 18 years)</p> <p>Diabetes (diagnosed before age 18 years)</p> <p>Emphysema</p> <p>Endometriosis</p> <p>Epilepsy/ Seizures</p> <p>Gout</p> <p>Headaches (>1 per week)</p> <p>Heart attack</p> <p>Heart valve problems</p> <p>Hepatitis</p> <p>Hypertension</p> <p>Hyperthyroid</p> <p>Hypothyroid</p> <p>Infections</p> <p>Multiple sclerosis</p> <p>Osteoporosis</p> <p>Other illnesses (please list)</p> <p>Rheumatoid arthritis</p> <p>Stomach/ Duodenal ulcers</p> <p>Stroke</p> <p>Tuberculosis</p> <p>Urinary incontinence</p> <p>Questionnaire contains 91 items divided into 8 sections (comorbidities, surgeries, medication, substance use and demographic characteristics)</p> | <p>100 preoperative orthopedic surgery patients at 1 hospital.</p> <p>Characteristics:</p> <p>Female 54%</p> <p>Mean age 46.9 years (\pm SD 16.7)</p> <p>College level or higher 64%</p> | <p>Comparison: NP/PA responses to identical questionnaire following medical record review and/or patient interview</p> <p>Results:</p> <p>Kappa: 0.15 (other illnesses) – 1.00; mean κ=0.69</p> <p>Sensitivity: -</p> <p>Specificity: -</p> <p>PPV: -</p> <p>NPV: -</p> | <p>Comparison: NP/PA responses to identical questionnaire following medical record review and/or patient interview</p> <p>Results:</p> <p>Mean percentage agreement across all questionnaire items between self-report and NP/PA report was 96%.</p> | - |
|----------------------------|--|---|---|--|---|

| | | | | | |
|---------------------|---|--|---|--|---|
| Fan ²⁸ | <p>Angina</p> <p>Arthritis</p> <p>CABG/ PTCA</p> <p>Cancer</p> <p>Congestive heart failure</p> <p>Coronary artery disease</p> <p>Depression</p> <p>Diabetes</p> <p>Drug abuse</p> <p>Enlarged prostate</p> <p>Heartburn</p> <p>HIV</p> <p>Hypertension</p> <p>Liver disease</p> <p>Lung disease</p> <p>Osteoporosis</p> <p>Pneumonia</p> <p>Post-Traumatic Stress Disorder</p> <p>Prior myocardial infarction</p> <p>Renal insufficiency</p> <p>Seizure</p> <p>Stroke</p> <p>Thyroid disease</p> <p>Ulcer disease</p> | <p>Development sample: 5469 patients from 7 VA medical centers. Data from the ACQUIP⁶⁵ study.</p> <p>Characteristics: Female 2.5% Mean age 67.8 years (\pm SD 0.1) Caucasian 83.4% College level or higher 68.7%</p> <p>Validation sample: 5478 patients from 7 VA medical centers</p> <p>Characteristics: Female 2.7% Mean age 67.8 years (\pm SD 0.1) Caucasian 83.3% College level or higher 68%</p> | - | - | <p>Comparison: -</p> <p>Results: Mortality: <i>All-cause mortality:</i> SIC had a moderate discriminative ability (AUC=0.71) of SIC in predicting mortality at 2 years follow-up. A combined model, containing SIC and SF-36 as predictors, had an AUC=0.74. PROM: - Health care utilization: <i>Re-hospitalizations:</i> Discriminative ability of SIC was less able in predicting 2-year re-hospitalizations (AUC=0.61), which slightly increased when SF-36 was added to the model (AUC=0.64).</p> |
| Lore ^{m27} | <p><i>All respondents:</i></p> <p>Angina</p> <p>Asthma</p> <p>Atopic eczema</p> <p>Cancer survivor</p> <p>Cerebrovascular stroke</p> <p>Chronic bronchitis</p> <p>Diabetes</p> <p>Duodenal ulcer</p> <p>Epilepsy</p> <p>Fibromyalgia</p> <p>Food allergies</p> <p>Hand eczema</p> <p>Hypersensitivity</p> <p>Kidney stone</p> <p>Liver disease</p> <p>Migraine</p> | <p>Reference population: 26684 patients sampled from Tromsø study (1994/1995)⁶⁶.</p> <p>Characteristics: Female 52.6% Age group <50 years 61.7%</p> <p>Validation population: 804 patients sampled from Tromsø study and FHI panel (2001/2002).</p> <p>Characteristics: Female 55% Ages 30-79 years</p> | - | <p>Comparison: Medical record-derived CCI</p> <p>Results: HII correlated more strongly with SRH vs. CCI (Spearman's $r=-0.360$, $p<0.001$ vs. $r=-0.250$, $p<0.001$). After excluding all patients with HII=0, the correlation between HII and SRH strengthened ($r=-0.421$, $p<0.001$) while it weakened between CCI and SRH ($r=-0.141$, $p<0.001$).</p> | <p>Comparison: -</p> <p>Results: Mortality: - PROM: <i>SRH:</i> In an ordinal logistic regression model (containing age, gender, mental health symptoms and the HII), HII had a negative effect ($\beta=-0.249$, $p<0.001$) after adjustment for the other variables. Health care utilization: -</p> |

| | | | | | |
|-------------------------------|--|---|---|---|---|
| | <p>Myocardial infarction Osteoporosis Pollen allergies Psoriasis Thyroid Ventricular ulcer</p> <p><i>For patients >70 years, added:</i> Arthritis Cataract Glaucoma Parkinson's disease Rheumatoid arthritis Urinary incontinence</p> | | | | |
| Luck e ³⁸ | <p><i>For analysis:</i> Asthma Cardiovascular disorder (combined) Coronary heart disease Diabetes mellitus Dyslipidemia GI Hypertension Hyperuricemia Mental disorders Osteoporosis</p> <p>Original questionnaire comprised of 51 (combined) diseases, as well as free text.</p> | <p>2653 patients with COPD or chronic bronchitis. Data from the COSYCONET⁶⁷ study.</p> <p>Characteristics: Male 59.4% Mean age 65 years (\pm SD 8.6) Mean BMI 27 (\pm SD 5.4) GOLD ≤ 2 57.7%</p> | <p>Comparison: ATC-codes for disease-specific medication</p> <p>Results: Concordance between self-reported comorbidities and ATC codes for disease-specific medication varied from 1.3% (asthma) to 51.8% (combined CVD).</p> | <p>Comparison: Matched ICD10-codes for diseases and non-specific medications</p> <p>Results: About 51.5% of self-reported comorbidities were confirmed after comparing them to matched ICD10-codes.</p> | - |
| Md Yus of ⁴⁴ | <p>18 sections: 12 physical systems 6 mental health state an additional free text option for medication prescription.</p> | <p>113 community-dwelling patients from 1 research center.</p> <p>Characteristics: Female 56.6% Mean age 75.3 years (\pm SD 5.19)</p> | - | <p>Comparison: Clinician report</p> <p>Results: CMI correlated significantly with the GP-data ($r=0.8$, $p<0.001$).</p> | <p>Comparison: -</p> <p>Results: Mortality: <i>Survival:</i> In a Cox proportional hazards model containing 5 continuous predictors (age, weighted CCI, combined condition and age-related CCI score, total score physical sections of CMI, total score mental sections of CMI, and count of medication prescriptions), none of</p> |

| | | | | | |
|-----------------------|---|--|---|--|--|
| | | | | | these predictors significantly contributed to predicting mortality. PROM: - Health care utilization: - |
| Merkin ³⁴ | Angioplasty or CABG Cancer Cerebrovascular disease Congestive Heart Failure COPD Diabetes Hypertension Myocardial infarction | 965 patients with ESRD from 81 dialysis clinics. Data from the CHOICE study ⁶⁸ . Characteristics: Female 46% Mean age 58 years Caucasian 67% College level or higher 36% | Comparison: Medical record-derived ICED Results: Kappa 0.19 (hypertension) – 0.93 (diabetes) Sensitivity: 18 (COPD) – 96% (diabetes) Specificity: 76 (hypertension) – 98% (diabetes) PPV: - NPV: - Comparison: Clinician report Results: Kappa 0.19 (hypertension) – 0.81 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: - | - | - |
| Mukerji ⁴⁵ | Arthritis Diabetes Heart disease Lung disease Other cancers (apart from index tumor) Psychiatric problems Stroke | 458 patients with newly diagnosed head-and-neck cancer from 3 hospitals. Characteristics: Female 23.6% Caucasian 86% College level or higher 49.8% | Comparison: Medical record-derived ACE-27 index Results: Kappa: 0.11 (arthritis) – 0.89 (diabetes) Sensitivity: - Specificity: - PPV: - NPV: - Over-reporting by self-report occurred more often (21.2%) compared to under-reporting | Comparison: Medical record-derived ACE-27 index Results: Kappa: 0.50 (0.44 – 0.57) | - |

| | | | | | |
|--------------------------|---|---|--|--|---|
| | | | (13.5%), with medical records as gold standard. | | |
| Paler j ⁶⁹ | 9 section headers: Heart & blood vessels Alcohol consumption Brain and nerves Cancer Diabetes Joints and muscles Kidney Liver, stomach and pancreas Lungs | 20 patients with head-and-neck cancer. Characteristics: - | - | Comparison: Medical record-derived ACE-27 grade Results: Kappa: 0.92 (95% CI 0.82 – 1.0) | - |
| Pati ² 5 | Acid peptic disease Arthritis Asthma Cancer Chronic back ache Chronic kidney disease Chronic liver disease (alcohol) Deafness Dementia Diabetes Epilepsy Filariasis Heart disease Hypertension Stroke Thyroid Tuberculosis Visual difficulty | 103 patients from 4 primary care practices. Characteristics: Female 45% Mean age 45 years (\pm SD 5.32) | Comparison: Clinician's prescription-derived data Results: Kappa: 0.58 (hearing problem) – 1.00 (tuberculosis); 16/18 conditions were evaluated. Sensitivity: - Specificity: - PPV: - NPV: - | - | - |
| Lee ² 9 | Alcohol use Arthritis BMI < 25 Chronic lung disease Current tobacco use Diabetes mellitus Heart failure History of falls History of pain Hypertension Incontinence | Older (>50 years) community-dwelling patients. Data from the Health and Retirement Study ^{70,71} . Development cohort: 11701 patients Characteristics: Female 57% | - | - | Results: - Mortality: <i>All-cause mortality</i> : a final model, predicting 4-year mortality, included 12 variables (6 comorbid conditions, sex, age, and 4 functional status measures). Discriminative ability of the model was determined in the development (ROC=0.84) and validation cohort (ROC=0.82). - PROM: - - Health care utilization: - |

| | | | | | |
|----------------------------|--|---|---|---|--|
| | Memory-related disease Non-skin cancers Other heart problems Psychiatric disease Stroke Visual or hearing impairment | Mean age 67 years (\pm SD 10) Caucasian 81% High school level or higher 75% Validation Cohort: 8009 patients Characteristics: Female 56% Mean age 67 years (\pm SD 10) Caucasian 71% High school level or higher 66% | | | |
| Voa klan der ³⁹ | Asthma Cancer Chronic back pain Circulatory problems Diabetes Digestive problems Emphysema Epilepsy Eye problems Hay fever/Allergies Heart disease Hypertension Kidney disease Liver disease Other Stomach ulcers Stroke Thyroid problems | 518 patients receiving major joint arthroplasty at 2 acute care facilities (1995-1997) 283 patients receiving TKA. Characteristics: Female 59% Mean age 69 years (\pm SD 9) 235 patients receiving THA. Characteristics: Female 60% Mean age 67.1 years (\pm SD 12) | - | Comparisons: 1. Medical record-derived CCI 2. Administrative data-derived CCI 3. ICD-9 Sum of comorbidities (available in the EHR) Results: SRC correlated significantly with the medical record-derived CCI (Spearman's $r=0.40$, $p<0.01$), administrative data-derived CCI (Spearman's $r=0.32$, $p<0.01$) and ICD-9 Sum (Spearman's $r=0.39$, $p<0.01$). | Comparisons: 1. Medical record-derived CCI 2. Administrative data-derived CCI 3. ICD-9 Sum of comorbidities Results: Mortality: - PROM: <i>SF-36 & WOMAC (health-related quality of life)</i> : SRC explained similar amounts in HRQoL domains compared to the ICD-9 Sum, medical record-derived CCI and administrative data-derived CCI. Health care utilization: <i>Hospital stay</i> : SRC explained the variance in acute length of stay less better compared to the other comorbidity measures. <i>ER visits</i> : SRC was slightly better in predicting emergency department visits than the other measures (2% variance). |
| Selim ²⁶ | <i>Physical conditions:</i> Angina pectoris Cancer Cataract Chronic low back pain Chronic lung disease | 2425 patients who previously received ambulatory care. Data from Veterans Health Study. Characteristics: | - | - | Comparison: - Results: Mortality: <i>Survival</i> : In a Cox proportional hazards model, the mortality risk was 14% for every increment |

| | | | | |
|--|---|--|--|---|
| <p> Congestive heart failure Diabetes Diverticulitis Enlarged prostate Gallbladder disease Gout Heart attack Hepatitis High blood pressure Inflammatory bowel disease Irregular heartbeat Osteoarthritis Osteoporosis Peptic ulcer Peripheral vascular disease Phlebitis Prostatitis Renal failure Rheumatoid arthritis Seizure Skin cancer Stroke Thyroid disease Transient ischemic attacks Urinary tract infection </p> <p> <i>Mental conditions:</i> Alcohol use Anxiety Bipolar Depression Posttraumatic stress disorder Schizophrenia </p> | <p> Female 0% Mean age 64 years (\pm SD 12.7) High school level or higher 41% </p> | | | <p> in physical CI which decreased to 9% ($p < 0.05$) after adjustment for sociodemographic and disability rating. PROM: <i>SF-36 (health-related quality of life):</i> Physical CI had the strongest correlations (Pearson's $r \geq 0.29$) with SF-36-PCS, while the mental CI correlated better with SF-36-MCS (Pearson's $r \geq 0.30$). Both physical and mental CI's were significantly correlated with all SF-36 scales. Combined physical/mental CI also correlated significantly with all HRQoL scales (Pearson's r range -0.29 – -0.45). Health care utilization: <i>Hospital visits:</i> Both physical and mental CI's had significant coefficients and explained 5% of the variance in total outpatient clinic visits, which increased to 7% after adjustment for sociodemographic and disability rating. </p> |
|--|---|--|--|---|

| | | | | | |
|--------------------|--|---|---|---|---|
| Vign ³¹ | Arthritis Cardiovascular disease Diabetes Gall bladder disease High cholesterol Hypertension Intestinal polyps Irritable bowel syndrome Osteoporosis Other cancers Thyroid disorders | 1936 breast cancer patients. Data from the SFBCS ⁷² & LACE ⁷³ studies. Characteristics: Mean age 60.4 years (\pm SD 11.0) Caucasian 71% College level or higher 67.8% | Comparison: Medical records (including diagnostic ICD-9 codes) Results: Kappa: 0.50 (other heart diseases) – 0.87 (diabetes) Sensitivity: 48 (other heart diseases) – 90.5% (myocardial infarction) Specificity: 96 (other heart diseases) – 99.5% (diabetes) PPV: - NPV: - | - | Comparison: Medical records (including diagnostic ICD-9 codes) Results: Mortality: <i>All-cause mortality:</i> No significant differences in HR's for covariates between comorbidity models with self-report vs. medical records as data source (diabetes HR 1.65 vs. 1.44; hypertension 1.22 vs. 1.55; myocardial infarction 1.40 vs. 1.73; other heart diseases 1.07 vs. 1.51) were observed. PROM: - Health care utilization: - |
|--------------------|--|---|---|---|---|

- = unknown/ not described in study; AIDS: Acquired Immuno-Deficiency Syndrome; SD: Standard Deviation; CCI: Charlson Comorbidity Index; PPV: positive predictive value; NPV: negative predictive value; r: (Spearman's) correlation coefficient; PROM: Patient-Reported Outcome Measure; ER: emergency room; RCT: randomized controlled trial; ICC: intra-class correlation coefficient; CI: confidence interval; ADL: Activities of Daily Living scale; AUC: area-under-the-curve; CTD: connective tissue disease; RA: rheumatoid arthritis; ASI: SCQ: Self-Administered Comorbidity Questionnaire; EQ-5D: EuroQol-5 Dimension; HER: electronic health record; SF-36-PCS: Short-Form 36 Physical Component Summary scale; SF-36-MCS: Short-Form 36 Mental Component Summary scale; β = beta; HIV: human immunodeficiency virus; COPD: chronic obstructive pulmonary disease; HR: hazard ratio; CHF: congestive heart failure; CAD: coronary artery disease; MMSE: Mini Mental Status Examination; mSCQ: modified version of Self-Administered Comorbidity Questionnaire; OASIS: Outcome in Ankylosing Spondylitis International Study; BASDAI: Bath Ankylosing Spondylitis Disease Activity Index; BASFI: Bath Ankylosing Spondylitis Functional Index; SCQ-G: German version of the Self-Administered Comorbidity Questionnaire; CORETH: The Choice of Renal Replacement Therapy project; HMO: Health Maintenance Organization; BRFSS: Behavior Risk Factor Surveillance System; GSE: General Self-Efficacy; DBMA-Fv: French version of the Disease Burden Morbidity Assessment; CIRS: Cumulative Illness Rating Scale; UTI: urinary tract infection; ELES-PS: Aging in Spain Longitudinal Study, Pilot Survey; PWI: Personal Wellbeing Index; CES-D: Center for Epidemiologic Studies Depression Scale; OA: osteo-arthritis; NEADL: Nottingham Extended Activities of Daily Living; CmSS: Comorbidity Symptom Scale; GHQ-28: General Health Questionnaire – 28; HAD: Hospital Anxiety and Depression Scale; CONNECT: Centralized Nurse-Led Telephone-Based Care Coordination to Improve Outcomes After Surgical Resection for Colorectal Cancer; PAOD: peripheral arterial occlusive disease; MS: multiple sclerosis; ICD-9: International Classification of Diseases version 9; IBD: inflammatory bowel disease; PCOS: Prostate Cancer Outcomes Study; NP: nurse practitioner; PA: physician assistant; PID: pelvic inflammatory disease; TIA: transient ischemic attack; CABG: coronary artery bypass graft; PTCA: percutaneous transluminal coronary angioplasty; VA: Veterans Affairs; ACQUIP: Ambulatory Care Quality Improvement Project; SIC: Seattle Index of Co-morbidity; FHI: Norwegian Institute of Public Health; HII: Health Impact Index; SRH: Self-Reported Health; COSYCONET: COPD and SYstemic consequences-COMorbidities NETwork cohort study; BMI: body mass index; ATC: Anatomical Therapeutic Chemical code; CVD: cardiovascular disorder; ICD-10: International Classification of Diseases version 10; CMI: Cornell Medical Index; GP: general practitioner; ESRD: end-stage renal disease; CHOICE: Choices for Healthy Outcomes in Caring for End-stage renal disease study; ICED: Index of Coexistent Disease; ACE-27: Adult Comorbidity Evaluation – 27; ROC: receiver-operating characteristic; TKA: total knee arthroplasty; THA: total hip arthroplasty; SRC: Self-report Co-Morbidity; WOMAC: Western Ontario McMaster Osteoarthritis Index; HRQoL: health-related quality-of-life; SFBCS: San Francisco Bay Area Breast Cancer Study; LACE: Life After Cancer Epidemiology study.